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The complete genome sequence of the Gram-positive bacterium *Bacillus subtilis*

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***Bacillus subtilis* is the best-characterized member of the Gram-positive bacteria. Its genome of 4,214,810 base pairs comprises 4,100 protein-coding genes. Of these protein-coding genes, 53% are represented once, while a quarter of the genome corresponds to several gene families that have been greatly expanded by gene duplication, the largest family containing 77 putative ATP-binding transport proteins. In addition, a large proportion of the genetic capacity is devoted to the utilization of a variety of carbon sources, including many plant-derived molecules. The identification of five signal peptidase genes, as well as several genes for components of the secretion apparatus, is important given the capacity of *Bacillus* strains to secrete large amounts of industrially important enzymes. Many of the genes are involved in the synthesis of secondary metabolites, including antibiotics, that are more typically associated with *Streptomyces* species. The genome contains at least ten prophages or remnants of prophages, indicating that bacteriophage infection has played an important evolutionary role in horizontal gene transfer, in particular in the propagation of bacterial pathogenesis.**

Techniques for large-scale DNA sequencing have brought about a revolution in our perception of genomes. Together with our understanding of intermediary metabolism, it is now realistic to envisage a time when it should be possible to provide an extensive chemical definition of many living organisms. During the past couple of years, the genome sequences of *Haemophilus influenzae*, *Mycoplasma genitalium*, *Synechocystis* PCC6803, *Methanococcus jannaschii*, *M. pneumoniae*, *Escherichia coli*, *Helicobacter pylori*, *Archaeoglobus fulgidus* and the yeast *Saccharomyces cerevisiae* have been published in their entirety¹⁻⁸, and at least 40 prokaryotic genomes are currently being sequenced. Regularly updated lists of genome sequencing projects are available at <http://www.mcs.anl.gov/home/gaasterl/genomes.html> (Argonne National Laboratory, Illinois, USA) and <http://www.tigr.org> (TIGR, Rockville, Maryland, USA).

The list of sequenced microorganisms does not currently include a paradigm for Gram-positive bacteria, which are known to be important for the environment, medicine and industry. *Bacillus subtilis* has been chosen to fill this gap^{9,10} as its biochemistry, physiology and genetics have been studied intensely for more than 40 years. *B. subtilis* is an aerobic, endospore-forming, rod-shaped bacterium commonly found in soil, water sources and in association with plants. *B. subtilis* and its close relatives are an important source of industrial enzymes (such as amylases and proteases), and much of the commercial interest in these bacteria arises from their capacity to secrete these enzymes at gram per litre concentrations. It has therefore been used for the study of protein secretion and for development as a host for the production of heterologous proteins¹¹. *B. subtilis* (*natto*) is also used in the production of Natto, a traditional Japanese dish of fermented soya beans.

Under conditions of nutritional starvation, *B. subtilis* stops growing and initiates responses to restore growth by increasing metabolic diversity. These responses include the induction of motility and chemotaxis, and the production of macromolecular hydrolases (proteases and carbohydrases) and antibiotics. If these responses fail to re-establish growth, the cells are induced to form chemically, irradiation- and desiccation-resistant endospores. Sporulation involves a perturbation of the normal cell cycle and the differentiation of a binucleate cell into two cell types. The division of the cell into a smaller forespore and a larger mother cell, each with an entire copy of the chromosome, is the first morphological indication of sporulation. The former is engulfed by the latter and differential expression of their respective genomes, coupled to a complex network of interconnected regulatory path-

ways and developmental checkpoints, culminates in the programmed death and lysis of the mother cell and release of the mature spore¹². In an alternative developmental process, *B. subtilis* is also able to differentiate into a physiological state, the competent state, that allows it to undergo genetic transformation¹³.

General features of the DNA sequence

Analysis at the replicon level. The *B. subtilis* chromosome has 4,214,810 base pairs (bp), with the origin of replication coinciding with the base numbering start point¹⁴, and the terminus at about 2,017 kilobases (kb)¹⁵. The average G + C ratio is 43.5%, but it varies considerably throughout the chromosome. This average is also different if one considers the nucleotide content of coding sequences, for which G and A (24% and 30%) are relatively more abundant than their counterparts C and T (20% and 26%). A significant inversion of the relative G - C/G + C ratio is visible at the origin of replication, indicating asymmetry of the nucleotide composition between the replication leading strand and the lagging strand¹⁶. Several A + T-rich islands are likely to reveal the signature of bacteriophage lysogens or other inserted elements (Fig. 1, see below).

We have analysed the abundance of oligonucleotides ('words') in the genome in various ways: absolute number of words in the genomic text, or comparison with the expected count derived from several models of the chromosome (for example, Markov models, or simulated sequences in which previously known features of the genome were conserved¹⁷). Comparing the experimental data with various models allowed us to define under- and overrepresentation of words in the experimental data set by reference to the model chosen. In general, the dinucleotide bias follows closely what has been described for other prokaryotes^{18,19}, in that the dinucleotides most overrepresented are AA, TT and GC, whereas those less represented are TA, AC and GT. Plots of the frequencies of AG, GA, CT and TC in sliding windows along the chromosome show dramatic decreases or increases around the origin and terminus of replication (data not shown). Trinucleotide frequency, directly related to the coding frame, will be discussed below. The distribution of words of four, five and six nucleotides shows significant correlations between the usage of some words and replication (several such oligonucleotides are very significantly overrepresented in one of the strands and underrepresented in the other one).

Setting a statistical cut-off for the significance of duplications at 10^{-3} , we expected duplication by chance of words longer than 24 nucleotides to be rare²⁰. In fact, the genome of *B. subtilis* contains a plethora of such duplications, some of them appearing more than

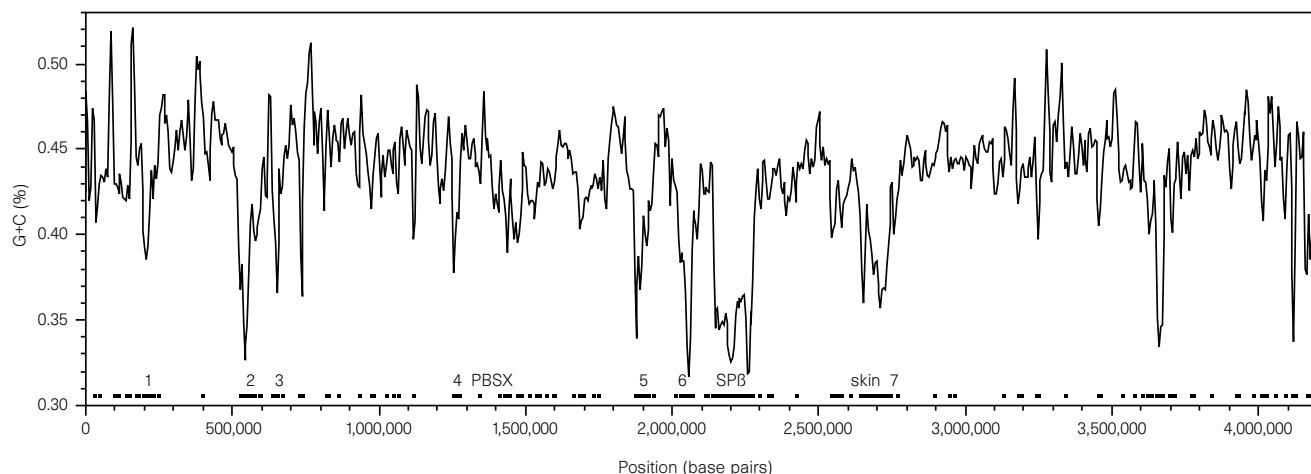


Figure 1 Distribution of A + T-rich islands along the chromosome of *B. subtilis*, in sliding windows of 10,000 nucleotides, with a step of 5,000 nucleotides. Location of genes from class 3 according to codon usage analysis (see Fig. 4) is indicated

by dots at the bottom of the graph. Known prophages (PBSX, SPB and *skin*) are indicated by their names, and prophage-like elements are numbered from 1 to 7.

twice. Among the duplications, we identified, as expected, the ribosomal RNA genes and their flanking regions, but also regions known to correspond to genes comprising long sequence repeats (such as *pks* and *srf*). We also found several regions that were not expected: a 182-bp repetition within the *yxaL* and *yxaO* genes; a 410-bp repetition between the *yxaK* and *yxaL* genes; an internal duplication of 174 bp inside *ycdI*; and significant duplications in the regions involved in the transcriptional control of several genes (such as 118 bp repeated three times between *yxbB* and *yxbC*). Finally, we found several repetitions at the borders of regions that might be involved in bacteriophage integration.

The most prominent duplication was a 190-bp element that was repeated 10 times in the chromosome. Multiple alignment of the ten repeats showed that they could be classified into two subfamilies with six and three copies each, plus a copy of what appears to be a chimera. Similar sequences have also been described in the closely related species *Bacillus licheniformis*^{21,22}. A striking feature of these repeats is that they are only found in half of the chromosome, at either side of the origin of replication, with five repeats on each side. Furthermore, with the exception of the most distal repeat at position 737,062, they lie in the same orientation with respect to the movement of the replication fork (Figs 2 and 3). Putative secondary structures conserved by compensatory mutations, as well as an insert in three of the copies, suggest that this element could indicate a structural RNA molecule.

Analysis at the transcription and translation level. Over 4,000 putative protein coding sequences (CDSs) have been identified, with an average size of 890 bp, covering 87% of the genome sequence (Fig. 2). We found that 78% of the genes started with ATG, 13% with TTG and 9% with GTG, which compares with 85%, 3% and 14%, respectively, in *E. coli*⁸. Fifteen genes (eight in the predicted CDSs in bacteriophage SP β) exhibiting unusual start codons (namely ATT and CTG) were also identified through their

similarities to known genes in other organisms or because they had a good GeneMark prediction (see Methods). This has not yet been substantiated experimentally. However, in the case of the gene coding for translation initiation factor 3, the similarity with its *E. coli* counterpart strongly suggests that the initiation codon is ATT, as is the case in *E. coli*.

We have not annotated CDSs that largely or entirely overlap existing genes, although such genes (for example, *comS* inside *srfAA*) certainly exist. It is also likely that some of the short CDSs present in the *B. subtilis* genome have been overlooked. For these reasons and possible sequencing errors, the estimated number of *B. subtilis* CDSs will fluctuate around the present figure of 4,100.

In several cases, in-frame termination codons or frameshifts were confirmed to be present on the chromosome (for example, an internal termination codon in *ywtF*, or the known programmed translational frameshift in *prfB*), indicating that the genes are either non-functional (pseudogenes) or subject to regulatory processes. It will therefore be of interest to determine whether these gene features are conserved in related *Bacillus* species, especially as strain 168 is derived from the Marburg strain that was subjected to X-ray irradiation²³.

A few regions do not have any identifiable feature indicating that they are transcribed: they could be 'grey holes' of the type described in *E. coli*²⁴. Preliminary studies involving all regions of more than 400 bp without annotated CDSs indicated that, of ~300 such regions, only 15% were likely to be really devoid of protein-coding sequences. One of the longest such regions, located between *yjfo* and *yjfn*, is 1,628 bp long. Grey holes seem generally to be clustered near the terminus of replication. However, a grey-hole cluster located at ~600 kb might be related to the temporary chromosome partition observed during the first stages of sporulation, when a segment of about one-third of the chromosome enters the prespore, and remains the sole part of the chromosome in the prespore for a significant transition period²⁵.

The codon usage of *B. subtilis* CDSs was analysed using factorial correspondence analysis¹⁷. We found that the CDSs of *B. subtilis* could be separated into three well-defined classes (Fig. 4). Class 1 comprises the majority of the *B. subtilis* genes (3,375 CDSs), including most of the genes involved in sporulation. Class 2 (188 CDSs) includes genes that are highly expressed under exponential growth conditions, such as genes encoding the transcription and translation machineries, core intermediary metabolism, stress proteins, and one-third of genes of unknown function. Class 3 (537 CDSs) contains a very high proportion of genes of unidentified function (84%), and the members of this class have codons enriched in A + T residues. These genes are usually clustered into groups between 15 and 160 genes (for example, bacteriophage SP β) and correspond to the A + T-rich islands described above (Fig. 1). When they are of known function, or when their products display similarity to proteins of known function, they usually correspond to functions found in, or associated with, bacteriophages or transposons, as well as functions related to the cell envelope. This includes the region *ycd/ydd/yde* (40 genes that are missing in some *B. subtilis* strains²⁶), where gene products showing similarities to bacteriophage and transposon proteins are intertwined. Many of these genes are associated with virulence genes identified in pathogenic Gram-positive bacteria, suggesting that such virulence factors are transmitted horizontally among bacteria at a much higher frequency than previously thought. If we include these A + T-rich regions as possible cryptic phages, together with known bacteriophages or bacteriophage-like elements (SP β , PBSX and the *skin* element), we find that the genome of *B. subtilis* 168 contains at least 10 such elements (Figs 2 and 3). Annotation of the corresponding regions often reveals the presence of genes that are similar to bacteriophage lytic enzymes, perhaps accounting for the observation that *B. subtilis* cultures are extremely prone to lysis.

The ribosomal RNA genes have been previously identified and

Table 1 Functional classification of the *Bacillus subtilis* protein-coding genes

The genes of known function or encoding products similar to known proteins in *B. subtilis* or in other organisms have been classified into functional categories (2,379 genes). The total number of genes in each category is indicated after the category title. Genes are listed in alphabetical order within each category, and their positions (in kilobases) on the *B. subtilis* chromosome are indicated after the gene names. A brief description is given for each gene. In some cases, interacting proteins have been indicated between brackets (for example, histidine kinases and response regulator, phosphatases and their substrates). More detailed and constantly updated information is available in the SubtiList database (see Methods). A preliminary assessment of the significance of sequence similarities was obtained through an automated procedure involving a combination between the BLAST2P probability and the percentage of amino-acid identity. Matches considered significant were re-examined manually. It should be emphasized that functions assigned to 'y' genes are based only on sequence similarity information with the best counterparts in protein databanks. Genes whose products are only similar to other unknown proteins, or not significantly similar to any other proteins in databanks (categories V and VI), were omitted.

Figure 2 General view of the *B. subtilis* chromosome. Arrows indicate the orientation of transcription. Genes are coloured according to their classification into six broad functional categories (blue, category I; green, category II; red, category III; orange, category IV; purple, category V; pink, category VI; see Table 1). Class 2 CDSs according to codon usage analysis are indicated by oblique hatches, and class 3 CDSs are indicated by vertical hatches. Ribosomal RNA genes are coloured in yellow. Transfer RNA genes are marked by triangles. Other RNA genes are represented as white arrows. Known genes (non-'y' genes) are printed in bold type. Putative transcription termination sites are represented as loops. Known prophages and prophage-like elements are indicated by brown hatches on the chromosome line. The 190-bp element repeated ten times is represented by hatched boxes.

shown to be organized into ten rRNA operons, mainly clustered around the origin of replication of the chromosome (Figs 2 and 3). In addition to the 84 previously identified tRNA genes, by using the Palingol²⁷ and tRNAscan²⁸ programs, we propose four putative new tRNA loci (at 1,262 kb, 1,945 kb, 2,003 kb and 2,899 kb), specific for lysine, proline and arginine (UUU, GGG, CCU and UCU anticodons, respectively). The 10S RNA involved in degradation of proteins made from truncated mRNA has been identified (*ssrA*), as well as the RNA component of RNase P (*rnpB*) and the 4.5S RNA involved in the secretion apparatus (*scr*).

There is a strong transcription orientation bias with respect to the movement of the replication fork: 75% of the predicted genes are transcribed in the direction of replication. Plotting the density of coding nucleotides in each strand along the chromosome readily identifies the replication origin and terminus (Fig. 3). To identify putative operons, we followed ref. 29 for describing Rho-independent transcription termination sites. This yielded ~1,630 putative terminators (340 of which were bidirectional). We retained only those that were located less than 100 bp downstream of a gene, or that were considered by the program to be 'very strong' (in order to account for possible erroneous CDSs). This yielded a total of ~1,250 terminators, with a mean operon size of three genes. A similar approach to the identification of promoters is problematical, especially because at least 14 sigma factors, recognizing different promoter sequences, have been identified in *B. subtilis*. Nevertheless, the consensus of the main vegetative sigma factor (σ^A) appears to be identical to its counterpart in *E. coli* (σ^{70}): 5'-TTGACA-*n*₁₇-TATAAT-3'. Relaxing the constraints of the similarity to sigma-specific consensus sequences led to an extremely high number of false-positive results, suggesting that the consensus-oriented approach to the identification of promoters should be replaced by another approach¹⁷.

Classification of gene products

Genes were classified according to ref. 14, based on the representation of cells as Turing machines in which one distinguishes between the machine and the program (Table 1). Using the BLAST2P software running against a composite protein databank compound of SWISS-PROT (release 34), TREMBL (release 3, update 1) and *B.*

subtilis proteins, we assigned at least one significant counterpart with a known function to 58% of the *B. subtilis* proteins. Thus for up to 42% of the gene products, the function cannot be predicted by similarity to proteins of known function: 4% of the proteins are similar only to other unknown proteins of *B. subtilis*; 12% are similar to unknown proteins from some other organism; and 26% of the proteins are not significantly similar to any other proteins in databanks. This preliminary analysis should be interpreted with caution, because only ~1,200 gene functions (30%) have been experimentally identified in *B. subtilis*. We used the 'y' prefix in gene names to emphasize that the function has not been ascertained (2,853 'y' genes, representing 70%).

Regulatory systems. Transcription regulatory proteins. Helix–turn–helix proteins form a large family of regulatory proteins found in both prokaryotes and eukaryotes. There are several classes, including repressors, activators and sigma factors. Using BLAST searches, we constructed consensus matrices for helix–turn–helix proteins to analyse the *B. subtilis* protein library. We identified 18 sigma or sigma-like factors, of which nine (including a new one) are of the SigA type. We also putatively identified 20 regulators (among which 18 were products of 'y' genes) of the GntR family, 19 regulators (15 'y' genes) of the LysR family, and 12 regulators (5 'y' genes) of the LacI family. Other transcription regulatory proteins were of the AraC family (11 members, 10 'y'), the Lrp family (7 members, 3 'y'), the DeoR family (6 members, 3 'y'), or additional families (such as the MarR, ArsR or TetR families). A puzzling observation is that several regulatory proteins display significant similarity to aminotransferases (seven such enzymes have been identified as showing similarity to repressors).

Two-component signal-transduction pathways. Two-component regulatory systems, consisting of a sensor protein kinase and a response regulator, are widespread among prokaryotes. We have identified 34 genes encoding response regulators in *B. subtilis*, most of which have adjacent genes encoding histidine kinases. Response regulators possess a well-conserved N-terminal phospho-acceptor domain³⁰, whereas their C-terminal DNA-binding domains share similarities with previously identified response regulators in *E. coli*, *Rhizobium meliloti*, *Klebsiella pneumoniae* or *Staphylococcus aureus*. Representatives of the four subfamilies recently identified in *E. coli*³¹

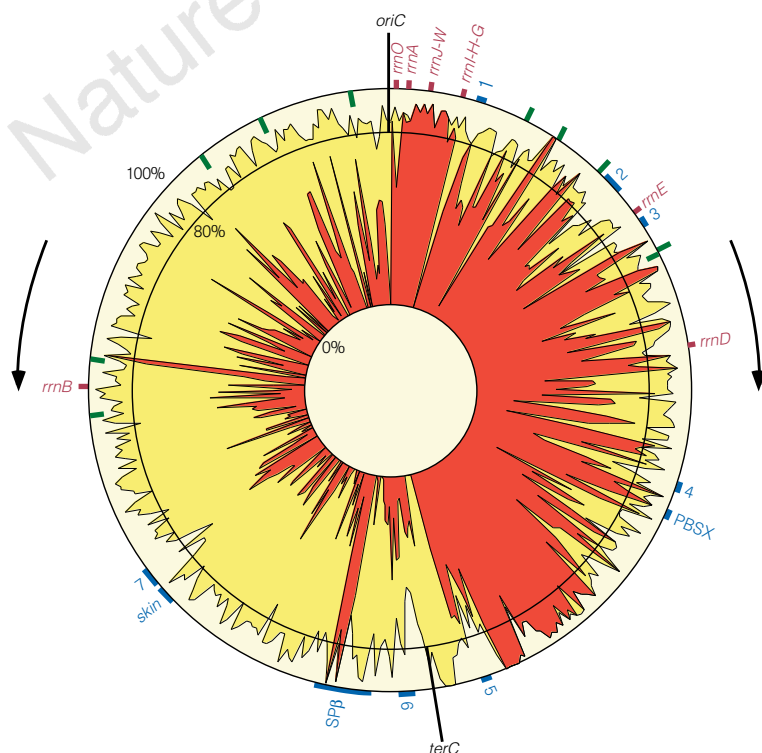


Figure 3 Density of coding nucleotides along the *B. subtilis* chromosome. Yellow stands for the density of coding nucleotides in both strands of the sequence; red indicates the density of coding nucleotides in the clockwise strand (nucleotides involved in genes transcribed in the clockwise orientation). The movement of the replication forks is represented by arrows. Ribosomal RNA operons are indicated by brown boxes. Known prophages and prophage-like elements are represented as blue lines. The 190-bp element repeated ten times is represented by green lines.

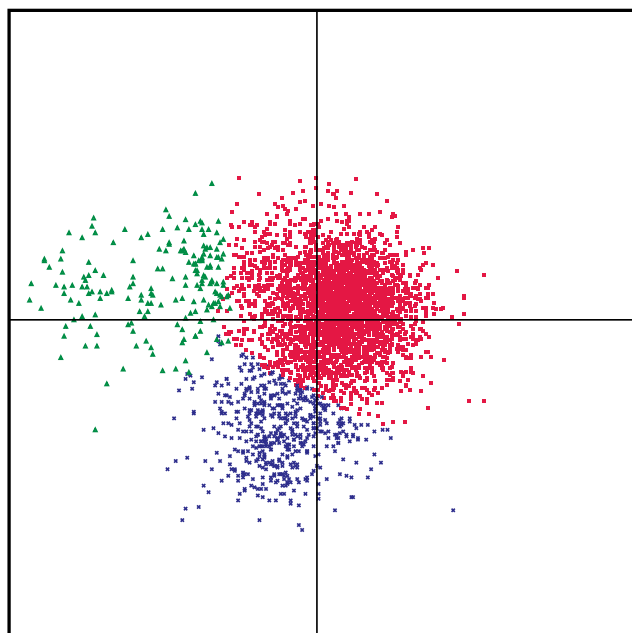


Figure 4 Factorial correspondence analysis of codon usage in the *B. subtilis* CDSs. Red dots, genes from class 1; green triangles, genes from class 2; blue crosses, genes from class 3. Class 2 contains genes coding for the translation and transcription machineries, and genes of the core intermediary metabolism. Class 3 genes correspond to codons strongly enriched in A or T in the wobble position; they generally belong to prophage-like inserts in the genome.

(OmpR, FixJ, CitB and LytR) have been identified in *B. subtilis*. In a fifth subfamily, CheY, the DNA-binding domain is absent. The DNA-binding domain of a single *B. subtilis* response regulator, YesN, shares similarity with regulatory proteins of the AraC family.

Quorum sensing. The *B. subtilis* genome contains 11 aspartate phosphatase genes, whose products are involved in dephosphorylation of response regulators, that do not seem to have counterparts in Gram-negative bacteria such as *E. coli*. Downstream from the corresponding genes are some small genes, called *phr*, encoding regulatory peptides that may serve as quorum sensors³². Seven *phr* genes have been identified so far, including three new genes (*phrG*, *phrI* and *phrK*).

Protein secretion. It is known that *B. subtilis* and related *Bacillus* species, in particular *B. licheniformis* and *B. amyloliquefaciens*, have a high capacity to secrete proteins into the culture medium. Several genes encoding proteins of the major secretion pathway have been identified: *secA*, *secD*, *secE*, *secF*, *secY*, *ffh* and *ftsY*. Surprisingly, there is no gene for the SecB chaperone. It is thought that other chaperone(s) and targeting factor(s), such as Ffh and FtsY, may take over the SecB function. Further, although there is only one such gene in *E. coli*, five type I signal peptidase genes (*sipS*, *sipT*, *sipU*, *sipV* and *sipW*) have been found³³. The *lsp* gene, encoding a type II signal peptidase required for processing of lipo-modified precursors, was also identified. PrsA, located at the outer side of the membrane, is important for the refolding of several mature proteins after their translocation through the membrane.

Other families of proteins. ABC transporters were the most frequent class of proteins found in *B. subtilis*. They must be extremely important in Gram-positive bacteria, because they have an envelope comprising a single membrane. ABC transporters will therefore allow such bacteria to escape the toxic action of many compounds. We propose that 77 such transporters are encoded in the genome. In general they involve the interaction of at least three gene products, specified by genes organized into an operon. Other families comprised 47 transport proteins similar to facilitators (and perhaps sometimes part of the ABC transport systems), 18 amino-acid permeases (probably antiporters), and at least 16 sugar transporters belonging to the PEP-dependent phosphotransferase system.

General stress proteins are important for the survival of bacteria under a variety of environmental conditions. We identified 43 temperature-shock and general stress proteins displaying strong similarity to *E. coli* counterparts.

Missing genes. Histone-like proteins such as HU and H-NS have been identified in *E. coli*. We found that *B. subtilis* encodes two putative histone-like proteins that show similarity to *E. coli* HU, namely HBSu and YonN, but found no homologue to H-NS. It is known that the *hbs* gene encoding HBSu is essential, but we do not expect the *yonN* gene to be essential because it is present in the SP β prophage. IHF is similar to HU, and it is not known whether HBSu plays a similar role to that of IHF in *E. coli*. Similarly, no protein similar to FIS could be found.

Genes encoding products that interact with methylated DNA, such as *seqA* in *E. coli*, involved in the regulation of replication initiation timing, or *mutH*, the endonuclease recognizing the newly synthesized strand during mismatch repair at hemi-methylated

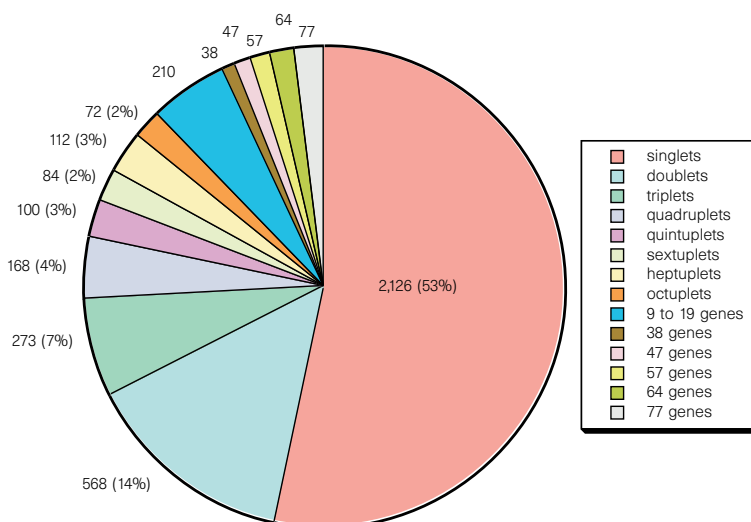


Figure 5 Gene paralogue distribution in the genome of *B. subtilis*. Each *B. subtilis* protein has been compared with all other proteins in the genome, using a Smith and Waterman algorithm. The baseline is established by making a similar

comparison using 100 independent random shuffles of the protein sequence (Z-score > 13).

GATC sites, are also missing. This is in line with the absence of known methylation in *B. subtilis*, equivalent to Dam methylation in *E. coli*. Similarly, *E. coli* *sfiA*, encoding an inhibitor of FtsZ action in the SOS response, has no counterpart in *B. subtilis*. In contrast, *B. subtilis* replication initiation-specific genes, such as *dnaB* and *dnaD*, are missing in *E. coli*. The exact counterpart of the *E. coli* *mukB* gene, involved in chromosome partitioning, does not exist in *B. subtilis*, but genes *spo0J* and *smc* (Smc is weakly similar to MukB), which are suggested to be involved in partitioning of the *B. subtilis* chromosome, are missing in *E. coli*.

Turnover of mRNA is controlled in *E. coli* by a 'degradosome' comprising RNase E. It has a counterpart in *B. subtilis*, but we failed to find a clear homologue of RNase E in this organism. Whether this is related to the role of ribosomal protein S1 as an RNA helicase involved in mRNA turnover in *E. coli* requires further investigation. In particular, a homologue of *rpsA* (S1 structural gene), *ypfD*, might be involved in a structure homologous to the degradosome³⁴.

Structurally unrelated genes of similar function. Several genes encode products that have similar functions in *E. coli* and *B. subtilis*, but have no evident common structure. This is the case for the helicase loader genes, *E. coli* *dnaC* and *B. subtilis* *dnaI*; the genes coding for the replication termination protein, *E. coli* *tus* and *B. subtilis* *rtp*; and the division topology specifier genes, *E. coli* *minE* and *B. subtilis* *divIVA*. The situation may even be more complex in multisubunit enzymes: *B. subtilis* synthesizes two DNA polymerase III α chains, one having 3'–5' proofreading exonuclease activity (PolC) and the other without the exonuclease activity (DnaE); in *E. coli*, only the latter exists. *E. coli* DNA polymerase II is structurally related to DNA polymerase α of eukaryotes, whereas *B. subtilis* YshC is related to DNA polymerase β .

Metabolism of small molecules

The type and range of metabolism used for the interconversion of low-molecular-weight compounds provide important clues to an organism's natural environment(s) and its biological activity. Here we briefly outline the main metabolic pathways of *B. subtilis* before the reconstruction of these pathways *in silico*, the correlation of genes with specific steps in the pathway, and ultimately the prediction of patterns of gene expression.

Intermediary metabolism. It has long been known that *B. subtilis* can use a variety of carbohydrates. As expected, it encodes an Embden–Meyerhof–Parnas glycolytic pathway, coupled to a functional tricarboxylic acid cycle. Further, *B. subtilis* is also able to grow anaerobically in the presence of nitrate as an electron acceptor. This metabolism is, at least in part, regulated by the FNR protein, binding to sites upstream of at least eight genes (four sites experimentally confirmed and four putative sites). A noteworthy feature of *B. subtilis* metabolism is an apparent requirement of branched short-chain carboxylic acids for lipid biosynthesis³⁵. Branched-chain 2-keto acid decarboxylase activity exists and may be linked to a variety of genes, suggesting that *B. subtilis* can synthesize and utilize linear branched short-chain carboxylic acids and alcohols.

Amino-acid and nucleotide metabolism. Pyrimidine metabolism of *B. subtilis* seems to be regulated in a way fundamentally different from that of *E. coli*, as it has two carbamylphosphate synthetases (one specific for arginine synthesis, the other for pyrimidine). Additionally, the aspartate transcarbamylase of *B. subtilis* does not act as an allosteric regulator as it does in *E. coli*. As in other microorganisms, pyrimidine deoxyribonucleotides are synthesized from ribonucleoside diphosphates, not triphosphates. The cytidine diphosphate required for DNA synthesis is derived from either the salvage pathway of mRNA turnover or from the synthesis of phospholipids and components of the cell wall. This means that polynucleotide phosphorylase is of fundamental importance in nucleic acid metabolism, and may account for its important role in competence³⁶. Two ribonucleoside reductases, both of class I, NrdEF type, are encoded by the *B. subtilis* chromosome, in one case

from within the SP β genome. In this latter case, the gene corresponding to the large subunit both contains an intron and codes for an intein (V.L., unpublished data). The gene of the small subunit of this enzyme also contains an intron, encoding an endonuclease, as was found for the homologue in bacteriophage T4.

By similarity with genes from other organisms, there appears to be, in addition to genes involved in amino-acid degradation (such as the *roc* operon, which degrades arginine and related amino acids), a large number of genes involved in the degradation of molecules such as opines and related molecules, derived from plants. This is also in line with the fact that *B. subtilis* degrades polygalacturonate, and suggests that, in its biotope, it forms specific relations with plants.

Secondary metabolism. In addition to many genes coding for degradative enzymes, almost 4% of the *B. subtilis* genome codes for large multifunctional enzymes (for example, the *srf*, *pps* and *pks* loci), similar to those involved in the synthesis of antibiotics in other genera of Gram-positive bacteria such as *Streptomyces*. Natural isolates of *B. subtilis* produce compounds with antibiotic activity, such as surfactin, fengycin and difficidin, that can be related to the above-mentioned loci. This bacterium therefore provides a simple and genetically amenable model in which to study the synthesis of antibiotics and its regulation. These pathways are often organized in very long operons (for example, the *pks* region spans 78.5 kb, about 2% of the genome). The corresponding sequences are mostly located near the terminus of replication, together with prophages and prophage-like sequences.

Paralogues and orthologues

It is important to relate intermediary metabolism to genome structure, function and evolution. We therefore compared the *B. subtilis* proteins with themselves, as well as with proteins from known complete genomes, using a consistent statistical method that allows the evaluation of unbiased probabilities of similarities between proteins^{37,38}. For *Z*-scores higher than 13, the number of proteins similar to each given protein does not vary, indicating that this cut-off value identifies sets of proteins that are significantly similar.

Families of paralogues. Many of the paralogues constitute large families of functionally related proteins, involved in the transport of compounds into and out of the cell, or involved in transcription regulation. Another part of the genome consists of gene doublets (568 genes), triplets (273 genes), quadruplets (168 genes) and quintuplets (100 genes). Finally, about half of the genome is made of genes coding for proteins with no apparent paralogues (Fig. 5). No large family comprises only proteins without any similarity to proteins of known function.

The process by which paralogues are generated is not well understood, but we might find clues by studying some of the duplications in the genome. Several approximate DNA repetitions, associated with very high levels of protein identity, were found, mainly within regions putatively or previously identified as prophages. This is in line with previous observations about PBSX and the *skin* element^{39,40}, and suggests that these prophage-like elements share a common ancestor and have diverged relatively recently. In addition, several protein duplications are in genes that are located very close to each other, such as *yukL* and *dhbF* (the corresponding proteins are 65% identical in an overlap of 580 amino acids), *yugJ* and *yugK* (proteins 73% identical), *yxjG* and *yxjH* (proteins 70% identical), and the entire *opuB* operon, which is duplicated 3 kb away (*opuC* operon, yielding ~80% of amino-acid identity in the corresponding proteins).

The study of paralogues showed that, as in other genomes, a few classes of genes have been highly expanded. This argues against the idea of the genome evolving through a series of duplications of ancestral genomes, but rather for the idea of genes as living organisms, subject to evolutionary constraints, some being sub-

mitted to expansion and natural selection, and others to local duplications of DNA regions.

Among paralogue doublets, some were unexpected, such as the three aminoacyl tRNA synthetases doublets (*hisS* (2,817 kb) and *hisZ* (3,588 kb); *thrS* (2,960 kb) and *thrZ* (3,855 kb); *tyrS* (3,036 kb) and *tyrZ* (3,945 kb)) or the two *mutS* paralogues (*mutS* and *yshD*). This latter situation is similar to that found in *Synechocystis*. In the case of *B. subtilis*, the presence of two MutS proteins could indicate that there are two different pathways for long-patch mismatch repair, possibly a consequence of the active genetic transformation mechanism of *B. subtilis*.

Families of orthologues. Because *Mycoplasma* spp. are thought to be derived from Gram-positive bacteria similar to *B. subtilis*, we compared the *B. subtilis* genome with that of *M. genitalium*. Among the 450 genes encoded by *M. genitalium*, the products of 300 are similar to proteins of *B. subtilis*. Among the 146 remaining gene products, a further 3 are similar to proteins of other *Bacillus* species, and 9 to proteins of other Gram-positive bacteria; 25 are similar to proteins of Gram-negative bacteria; and 19 are similar to proteins of other *Mycoplasma* spp. This leaves only 90 genes that would be specific to *M. genitalium* and might be involved in the interaction of this organism with its host.

The *B. subtilis* genome is similar in size to that of *E. coli*. Because these bacteria probably diverged more than one billion years ago, it is of evolutionary value to investigate their relative similarity. About 1,000 *B. subtilis* genes have clear orthologous counterparts in *E. coli* (one-quarter of the genome). These genes did not belong either to the prophage-like regions or to regions coding for secondary metabolism (~15% of the *B. subtilis* genome). This indicates that a large fraction of these genomes shared similar functions. At first sight, however, it seems that little of the operon structure has been conserved. We nevertheless found that ~100 putative operons or parts of operons were conserved between *E. coli* and *B. subtilis*. Among these, ~12 exhibited a reshuffled gene order (typically, the arabinose operon is *araABD* in *B. subtilis* and *araBAD* in *E. coli*). In addition to the core of the translation and transcription machinery, we identified other classes of operons that were well conserved between the two organisms, including major integrated functions such as ATP synthesis (*atp* operon) and electron transfer (*cta* and *qox* operons). As well as being well preserved, the murein biosynthetic region was partly duplicated, allowing creation of part of the genes required for the sporulation division machinery⁴¹. The amino-acid biosynthesis genes differ more in their organization: the *E. coli* genes for arginine biosynthesis are spread throughout the chromosome, whereas the arginine biosynthesis genes of *B. subtilis* form an operon. The same is true for purine biosynthetic genes. Genes responsible for the biosynthesis of coenzymes and prosthetic groups in *B. subtilis* are often clustered in operons that differ from those found in *E. coli*. Finally, several operons conserved in *E. coli* and *B. subtilis* correspond to unknown functions, and should therefore be priority targets for the functional analysis of these model genomes.

Comparison with *Synechocystis* PCC6803 revealed about 800 orthologues. However, in this case the putative operon structure is extremely poorly conserved, apart from four of the ribosomal protein operons, the *groES*–*groEL* operon, *yfnHG* (respectively in *Synechocystis* *rfbFG*), *rpsB-ts*, *ylxS-nusA-infB*, *asd-dapGA-ymfA*, *spmAB*, *efp-accB*, *grpE-dnaK*, *yurXW*. The nine-gene *atp* operon of *B. subtilis* is split into two parts in *Synechocystis*: *atpBE* and *atpIHGFDAC*.

Conclusion

The biochemistry, physiology and molecular biology of *B. subtilis* have been extensively studied over the past 40 years. In particular, *B. subtilis* has been used to study postexponential phase phenomena such as sporulation and competence for DNA uptake. The genome sequences of *E. coli* and *B. subtilis* provide a means of studying the

evolutionary divergence, one billion years ago, of eubacteria into the Gram-positive and Gram-negative groups. The availability of powerful genetic tools will allow the *B. subtilis* genome sequence data to be exploited fully within the framework of a systematic functional analysis program, undertaken by a consortium of 19 European and 7 Japanese laboratories coordinated by S. D. Ehrlich (INRA, Jouy-en-Josas, France) and by N. Ogasawara and H. Yoshikawa (Nara Institute of Science and Technology, Nara, Japan). □

Methods

Genome cloning and sequencing. An international consortium was established to sequence the genome of *B. subtilis* strain 168 (refs 9, 10, 42). At its peak, 25 European, seven Japanese and one Korean laboratory participated in the program, together with two biotechnology companies. Five contiguous DNA regions totalling 0.94 Mb, and two additional regions of 0.28 and 0.14 Mb, were sequenced by the Japanese partners, while the European partners sequenced a total of 2.68 Mb. A few sequences from strain 168 published previously were not resequenced when long overlaps did not indicate differences.

A major technical difficulty was the inability to construct in *E. coli* gene banks representative of the entire *B. subtilis* chromosome using vectors that have proved efficient for other sources of bacterial DNA (such as bacteriophage or cosmid vectors). This was due to the generally very high level of expression of *B. subtilis* genes in *E. coli*, leading to toxic effects. This limitation was overcome by: cloning into a variety of vectors^{9,43,44}; using an *E. coli* strain maintaining low-copy number plasmids⁴⁴; using an integrative plasmid/marker rescue genome-walking strategy⁴⁴; and *in vitro* amplification using polymerase chain reaction (PCR) techniques^{45,46}.

Although cloning vectors were used in the early stages as templates for sequencing reactions, they were largely superseded in the later stages by long-range and inverse PCR techniques. To reduce sequencing errors resulting from PCR amplification artefacts, at least eight amplification reactions were performed independently and subsequently pooled. The various sequencing groups were free to choose their own strategy, except that all DNA sequences had to be determined entirely on both strands.

Sequence annotation and verification. The sequences were annotated by the groups, and sent to a central depository at the Institut Pasteur¹⁴. The Japanese sequences were also sent there through the Japanese depository at the Nara Institute of Science and Technology. The same procedures were used to identify CDSs and to detect frameshifts. They were embedded within a cooperative computer environment dedicated to automatic sequence annotation and analysis³⁹. In a first step, we identified in all six possible frames the open reading frames (ORFs) that were at least 100 codons in length. In a second step, three independent methods were used: the first method used the GeneMark coding-sequence prediction method⁴⁷ together with the search for CDSs preceded by typical translation initiation signals (5'-AAGGAGGTG-3'), located 4–13 bases upstream of the putative start codons (ATG, TTG or GTG); the second method used the results of a BLAST2X analysis performed on the entire *B. subtilis* genome against the non-redundant protein database at the NCBI; and the third method was based on the distribution of non-overlapping trinucleotides or hexanucleotides in the three frames of an ORF⁴⁸.

In general, frameshifts and missense mutations generating termination codons or eliminating start codons are relatively easy to detect. We shall devise a procedure for detecting another type of error, GC instead of CG or vice versa, which are much more difficult to identify. It should be noted that putative frameshift errors should not be corrected automatically. The sequences of the flanking regions of a 500-bp fragment centred around a putative error were sent to an independent verification group, which performed PCR amplifications using chromosomal DNA as template, and sequenced the corresponding DNA products.

Organization and accessibility of data. The *B. subtilis* sequence data have been combined with data from other sources (biochemical, physiological and genetic) in a specialized database, SubtiList⁴⁹, available as a Macintosh or Windows stand-alone application (4th Dimension runtime) by anonymous FTP at <ftp://ftp.pasteur.fr/pub/GenomeDB/SubtiList>. SubtiList is also accessible through a World-Wide Web server at <http://www.pasteur.fr/Bio/SubtiList.html>,

where it has been implemented on a UNIX system using the Sybase relational database management system. A completely rewritten version of SubtiList is in preparation to facilitate browsing of the information of the whole chromosome. Flat files of the whole DNA and protein sequences in EMBL and FASTA format will be made available at the above ftp address. Another *B. subtilis* genome database is also under development at the Human Genome Center of Tokyo University (<http://www.genome.ad.jp>), and SubtiList will also be available there.

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Table 1. Functional classification of the *Bacillus subtilis* protein-coding genes.

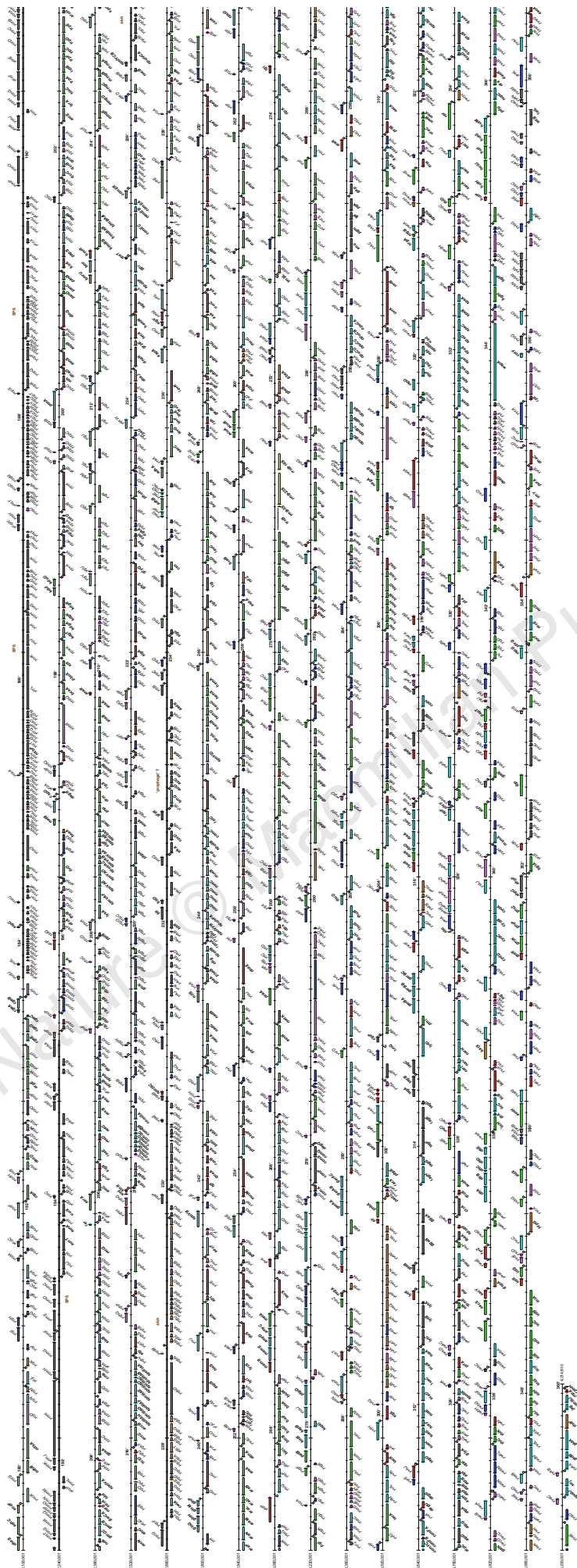
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ydiA	578	antibiotic resistance protein	ytmI	3007	amino acid ABC transporter (binding protein)	yciK	427	two-component sensor histidine kinase [YciU]
ydiI	580	arsenical pump membrane protein	ytmK	3006	amino acid ABC transporter (binding protein)	ydbF	497	two-component sensor histidine kinase [YdbG]
ydiJ	589	antibiotic transport-associated protein	ytmL	3006	amino acid ABC transporter (permease)	ydhH	587	two-component sensor histidine kinase [YdhI]
ydiU	590	multidrug-efflux transporter regulator	ytmM	3005	amino acid ABC transporter (permease)	yeshM	590	two-component sensor histidine kinase [YeshN]
ydiA	596	cation efflux system	ytmB	3125	aniline permease	ytiI	903	two-component sensor histidine kinase [YtiK]
ydiO	597	ABC transporter (binding protein)	ytmB	3118	ABC transporter (ATP-binding protein)	yhcY	1008	two-component sensor histidine kinase [YhcZ]
ydgF	608	amino acid ABC transporter (permease)	ytrE	3115	ABC transporter (ATP-binding protein)	yhlL	1129	sensory transduction pleiotropic regulatory protein
ydgH	609	transporter	ytsC	3111	ABC transporter (ATP-binding protein)			
ydgK	613	bicyclomycin resistance protein	ytsD	3110	ABC transporter (permease)	ykoH	1392	two-component sensor histidine kinase [YkoG]
ydhL	626	chloramphenicol resistance protein	yttB	3108	multidrug resistance protein	ykrQ	1419	two-component sensor histidine kinase
ydhM	626	cellulose phosphotransferase system enzyme II	yudB	3192	multidrug resistance protein	ykcD	1432	two-component sensor histidine kinase
ydhN	627	cellulose phosphotransferase system enzyme II	yugG	3188	Na ⁺ -transporting ATP synthase	yocV	2090	two-component sensor histidine kinase [YocG]
ydhO	627	cellulose phosphotransferase system enzyme II	yuiO	3239	ABC transporter (lipoprotein)	yrrP	2704	two-component sensor histidine kinase [YrrP]
ydiF	646	ABC transporter (ATP-binding protein)	yuiO	3240	ABC transporter (ATP-binding protein)	ytrP	3035	two-component sensor histidine kinase
ydiF	658	H ⁺ -symporter	yurF	3244	organic acid transport protein	ytsB	3112	two-component sensor histidine kinase [YtsA]
ydiJ	674	sugar transporter	yurJ	3248	Na ⁺ /H ⁺ antiporter	yurL	3566	two-component sensor histidine kinase [YurM]
ydiB	687	cation efflux system membrane protein	yurV	3249	Na ⁺ /H ⁺ antiporter	yvcQ	3566	two-component sensor histidine kinase [YvcP]
yecA	712	amino acid permease	yugQ	3218	potassium channel protein	yvtT	3497	two-component sensor histidine kinase [YvtU]
yecO	761	sugar-binding protein	yuniI	3330	purine permease	yvqB	3385	two-component sensor histidine kinase [YvqA]
yepP	762	lactose permease	yuniK	3331	purine permease	yvqE	3395	two-component sensor histidine kinase [YvqG]
yepS	763	lactose permease	yurJ	3345	multiple sugar ABC transporter (ATP-binding protein)	yvrG	3407	two-component sensor histidine kinase [YvrH]
yfhA	921	iron(III) dicitrate transport permease				ywdD	3741	two-component sensor histidine kinase
yfhI	926	antibiotic resistance protein	yurM	3348	sugar permease	yxdK	4071	two-component sensor histidine kinase [YxdL]
yfhB	933	ABC transporter (ATP-binding protein)	yurN	3349	sugar permease	yxiM	3992	two-component sensor histidine kinase [YxiL]
yfiC	896	ABC transporter (ATP-binding protein)	yurO	3350	multiple sugar-binding protein	yycG	4153	two-component sensor histidine kinase [YycF]
yfiG	900	metabolite transport protein	yurY	3360	ABC transporter (ATP-binding protein)			
yfiS	905	ABC transporter (ATP-binding protein)	yurY	3363	ABC transporter (ATP-binding protein)	I.4	MEMBRANE BIOENERGETICS (ELECTRON	
yfiM	907	ABC transporter (ATP-binding protein)	yusP	3374	multidrug-efflux transporter	SYNTHASE)	78	
yfiN	907	ABC transporter (ATP-binding protein)	yusV	3379	iron(III) dicitrate transport permease			
yfiS	913	multidrug resistance protein	yurK	3307	Na ⁺ /nucleoside cotransporter	atpA	3784	ATP synthase (subunit α)
yfiU	916	multidrug-efflux transporter	yuxJ	3232	multidrug-efflux transporter	atpB	3787	ATP synthase (subunit α)
yfiY	920	iron(III) dicitrate transport permease	yvaE	3448	multidrug-efflux transporter	atpC	3781	ATP synthase (subunit ϵ)
yfiZ	920	iron(III) dicitrate transport permease	yvbW	3490	amino acid permease	atpD	3782	ATP synthase (subunit β)
yfiQ	872	divalent cation transport protein	yvcC	3579	ABC transporter (ATP-binding protein)	atpE	3786	ATP synthase (subunit ϵ)
ykfE	865	H ⁺ /Ca ²⁺ exchanger	yvcR	3565	ABC transporter (ATP-binding protein)	atpF	3786	ATP synthase (subunit β)
ykfF	866	multidrug-efflux transporter	yvcS	3565	ABC transporter (permease)	atpG	3783	ATP synthase (subunit γ)
ykfH	862	transporter	yvdB	3561	transporter	atpH	3785	ATP synthase (subunit δ)
ykfL	861	multidrug resistance protein	yvdG	3555	maltose/maltodextrin-binding protein	atpI	3787	ATP synthase (subunit ι)
yfiA	844	amino acid carrier protein	yveA	3554	maltodextrin transport system permease	cccA	2599	cytochrome c_{550}
yfiE	844	anion-binding protein	yvcl	3552	maltodextrin transport system permease	ccbB	3625	cytochrome c_{551}
yfiF	840	phosphotransferase system enzyme II	yveA	3538	permease	ccaA	1922	required for a late step of cytochrome c synthesis
yfiS	829	2-oxoglutarate/malate translocator	yvfiH	3510	L-lactate permease	ctaA	1558	cytochrome caa_3 oxidase (required for biosynthesis)
yfmD	826	ferrichrome ABC transporter (binding protein)	yvkK	3508	maltose/maltodextrin-binding protein	ctaB	1559	cytochrome caa_3 oxidase (assembly factor)
yfmD	825	ferrichrome ABC transporter (permease)	yvfiL	3506	maltodextrin transport system permease	ctaC	1560	cytochrome caa_3 oxidase (subunit II)
yfmE	824	ferrichrome ABC transporter (permease)	yvfiM	3505	maltodextrin transport system permease	ctaD	1561	cytochrome caa_3 oxidase (subunit I)
yfmF	823	ferrichrome ABC transporter (ATP-binding protein)	yvfiR	3498	ABC transporter (ATP-binding protein)	ctaE	1563	cytochrome caa_3 oxidase (subunit III)
yfmM	815	ABC transporter (ATP-binding protein)	yvfiK	3424	molybdenum-binding protein	ctaF	1563	cytochrome caa_3 oxidase (subunit IV)
yfmO	812	multidrug-efflux transporter	yvfiL	3424	molybdate-binding protein	cydA	3978	cytochrome bd ubiquinol oxidase (subunit I)
yfmR	809	ABC transporter (ATP-binding protein)	yvfiM	3425	molybdenum transport permease	cydB	3977	cytochrome bd ubiquinol oxidase (subunit II)
yfmA	806	metabolite transport protein	yvfiW	3440	heavy metal-transporting ATPase	eifA	2915	electron transfer flavoprotein (α subunit)
ygaD	939	ABC transporter (ATP-binding protein)	yvfiX	3443	mercuric transport protein	eifB	2916	electron transfer flavoprotein (β subunit)
ygaL	961	nitrate ABC transporter (binding protein)	yvfiX	3443	mercuric transport protein	fer	2409	ferredoxin
ygaM	963	ABC transporter (permease)	yvfiA	3618	multidrug-efflux transporter	hmp	1372	flavohemoglobin
ygaB	962	ABC transporter (binding lipoprotein)	yvfiA	3605	transporter	narG	3829	nitrate reductase (α subunit)
yhaQ	1062	ABC transporter (ATP-binding protein)	yvfiJ	3399	macrolide-efflux protein	narH	3825	nitrate reductase (β subunit)
yhaU	1060	Na ⁺ /H ⁺ antiporter	yvfiJ	3402	iron transport system	narI	3823	nitrate reductase (γ subunit)
yhcA	977	multidrug resistance protein	yvfiC	3403	iron-binding protein	narJ	3824	nitrate reductase (protein I)
yhcG	981	glycine betaine/L-proline transport	yvfiC	3403	iron-binding protein	ndhF	205	NADH dehydrogenase (subunit 5)
yhcH	982	ABC transporter (ATP-binding protein)	yvfiO	3413	amino acid ABC transporter (ATP-binding protein)	qcrA	2364	menaquinone:cytochrome c oxidoreductase (iron-sulphur subunit)
yhcI	984	ABC transporter (binding lipoprotein)	yvfiH	3420	ABC transporter (amino acid permease)			
yhcL	986	sodium-glutamate symporter	yvfiA	3338	phosphotransferase system enzyme II	qcrB	2364	menaquinone:cytochrome c oxidoreductase (cytochrome b subunit)
yhdC	1023	amino acid transporter	yvfiF	3333	Na ⁺ -dependent symport	qcrC	2363	menaquinone:cytochrome c oxidoreductase (cytochrome b/c subunit)
yhdH	1024	sodium-dependent transporter	yvfiF	3333	Na ⁺ -dependent symport			
yheH	1047	ABC transporter (ATP-binding protein)	yvfiC	3904	nitrite transporter	qoxA	3917	cytochrome aa_3 quinol oxidase (subunit II)
yheI	1045	ABC transporter (ATP-binding protein)	yvfiA	3874	chloramphenicol resistance	qoxB	3916	cytochrome aa_3 quinol oxidase (subunit I)
yheL	1044	Na ⁺ /H ⁺ antiporter	yvfiF	3869	efflux protein	qoxC	3914	cytochrome aa_3 quinol oxidase (subunit III)
yhiQ	1107	iron(III) dicitrate-binding protein	yvfiQ	3837	ABC transporter (ATP-binding protein)	qoxD	3913	cytochrome aa_3 quinol oxidase (subunit IV)
yhiB	1120	metabolite permease	yvfiA	3821	ABC transporter (ATP-binding protein)	resA	2421	essential protein similar to cytochrome c biogenesis protein
yhiO	1133	multidrug-efflux transporter	yvoA	3758	bacteriocin transport permease			
yhiP	1133	transporter binding protein	ywoD	3754	transporter	resB	2420	essential protein similar to cytochrome c biogenesis protein
yhiG	1177	multidrug resistance protein	ywoE	3753	permease	resC	2418	essential protein similar to cytochrome c biogenesis protein
yizZ	1194	multidrug resistance protein	ywoG	3749	antibiotic resistance protein	tlp	1930	thioredoxin-like protein
yjbQ	1240	Na ⁺ /H ⁺ antiporter	ywcF	3743	large conductance mechanosensitive channel	trxA	2912	thioredoxin
yjdA	1272	fructose phosphotransferase system enzyme II	ywvA	3721	chromate transport protein	trxB	2912	thioredoxin
yjkb	1206	amino acid ABC transporter (ATP-binding protein)	ywvB	3720	chromate transport protein	trxB	2912	thioredoxin
yjmb	1301	Na ⁺ /galactoside symporter	ywvB	3720	chromate transport protein	trxB	2912	thioredoxin
yjmg	1307	hexuronate transporter	ywvK	3712	arsenical pump membrane protein	trxB	2912	thioredoxin
ykaB	1350	low-affinity inorganic phosphate transporter	ywvG	3693	metabolite transport protein	trxB	2912	thioredoxin
ykaB	1352	amino acid permease	yxaM	4100	antibiotic resistance protein	ycgT	352	thioredoxin reductase
ykcA	1353	ABC transporter (binding protein)	yxcC	4087	metabolite transport protein	ygdD	439	NAD(P)-flavin oxidoreductase
yktD	1368	oligopeptide ABC transporter (permease)	yxdL	4070	ABC transporter (ATP-binding protein)	ydbP	508	thioredoxin
yknU	1499	ABC transporter (ATP-binding protein)	yxdM	4069	ABC transporter (permease)	ydeQ	576	NAD(P)H oxidoreductase
yknV	1501	ABC transporter (ATP-binding protein)	yxeB	4066	ABC transporter (binding protein)	ydfQ	598	thioredoxin
ykoD	1390	cation ABC transporter (ATP-binding protein)	yxeM	4059	amino acid ABC transporter (binding protein)	ydgI	613	NADH dehydrogenase
yknS	1395	amino acid ABC transporter (permease)	yxeN	4058	amino acid ABC transporter (permease)	ykoO	814	NAD(P)-flavin oxidoreductase
ykpA	1512	ABC transporter (ATP-binding protein)	yxeR	4054	amino acid ABC transporter (ATP-binding protein)	yfnI	818	quinone oxidoreductase
ykrM	1416	Na ⁺ -transporting ATP synthase	yxiQ	4009	Mg ²⁺ /citrate complex transporter	yjcd	1280	cytochrome c oxidase assembly factor
ykuC	1476	macrolide-efflux protein	yxiQ	4009	Mg ²⁺ /citrate complex transporter	yjcd	1280	cytochrome c oxidase assembly factor
ykvW	1451	heavy metal-transporting ATPase	yxiJ	4005	pyrimidine nucleoside transport	ykuN	1486	flavodoxin
ymlA	1606	ABC transporter (ATP-binding protein)	yxiJ	3979	metabolite-sodium symport	ykuP	1488	sulfite reductase
ylnA	1630	anion permease	yxiA	3970	purine/cytosine permease	ykuU	1492	2-cys peroxiredoxin
yloB	1637	calcium-transporting ATPase	yxiF	3968	ABC transporter (ATP-binding protein)	ykvV	1450	thioredoxin
yml	1887	H ⁺ -symporter	yxiH	3966	multidrug-efflux transporter	yneN	1929	thiol:disulfide interchange protein
yncC	1896	metabolite transport protein	yylA	4194	transporter	yoiJN	2114	nitric-oxide reductase
yocN	2038	permease	yylB	4180	antibiotic resistance protein	yoiL	2287	thioredoxin
yocR	2106	sodium-dependent transporter	yylB	4175	ABC transporter (ATP-binding protein)	yosR	2159	thioredoxin
yocS	2109	sodium-dependent transporter	yylB	4174	ABC transporter (permease)	ypdA	2401	thioredoxin reductase
yocD	2129	aromatic metabolite transporter	yylB	4169	ABC transporter (permease)	yqiG	2516	NAD(P)-dependent flavin oxidoreductase
yodF	2130	proline permease	yycB	4159	ABC transporter (permease)	yqmM	2475	NAD(P)-dependent flavin oxidoreductase
yojA	2125	gluconate permease	yylD	4125	ABC transporter (ATP-binding protein)	yrlK	2708	NAD(P)H oxidoreductase
yqpE	2337	phosphotransferase system enzyme II	yzeE	4122	phosphotransferase system enzyme II	ythA	3139	cytochrome d oxidase subunit
yqgW	2620	Na ⁺ /P _i cotransporter				ytpP	3054	thioredoxin H1
yqgG	2581	phosphate ABC transporter (binding protein)	I.3	SENSORS (SIGNAL TRANSDUCTION)	38	ytrC	3117	cytochrome c oxidase subunit
yqgH	2580	phosphate ABC transporter (permease)	cheA	1712	two-component sensor histidine kinase [CheB/CheY] chemotactic signal modulator	ytrD	3116	cytochrome c oxidase subunit
yqgl	2579	phosphate ABC transporter (permease)	citS	830	two-component sensor histidine kinase [CitT]	yuiD	3249	NADH dehydrogenase (ubiquinone)
yqgl	2578	phosphate ABC transporter (ATP-binding protein)	compP	3255	two-component sensor histidine kinase [ComA]	yuiT	3246	NADH dehydrogenase
yqgK	2577	phosphate ABC transporter (ATP-binding protein)				yumb	3300	NADH dehydrogenase
yqgK	2575	lipoprotein	degS	3646	two-component sensor histidine kinase [DegU]	yumC	3301	thioredoxin reductase
yqiK	2492	amino acid ABC transporter (binding protein)				yusE	3354	thioredoxin
yqiY	2491	amino acid ABC transporter (permease)				yusF	3308	NADH dehydrogenase
yqiZ	2491	amino acid ABC transporter (ATP-binding protein)	kinA	1469	two-component sensor histidine kinase [SpoOF]	yvaB	3445	NAD(P)H dehydrogenase (quinone)
yqiV	2466	multidrug resistance protein				ywcG	3911	NAD(P)-flavin oxidoreductase
yqkI	2453	Na ⁺ /H ⁺ antiporter	kinB	3229	two-component sensor histidine kinase [SpoOF]	ywhN	3840	ubiquinol-cytochrome c reductase
yraO	2745	citrate transporter				ywrO	3708	NAD(P)H oxidoreductase
yrbD	2841	sodium/proton-dependent alanine carrier protein	kinC	1518	two-component sensor histidine kinase [SpoOA]			
ybdD	2968	antibiotic resistance protein				I.5	MOBILITY AND CHEMOTAXIS	65
ytcP	3087	ABC transporter (permease)				cheC	1715	inhibition of CheR-mediated methylation of methyl-accepting chemotaxis proteins
ytcQ	3086	lipoprotein	lysT	2957	two-component sensor histidine kinase [LysT]	cheD	1715	required for methylation of methyl-accepting chemotaxis proteins by CheR
ytcQ	3082	sugar transport protein				cheR	2380	methyl-accepting chemotaxis proteins methyltransferase
ytcQ	3145	ABC transporter (membrane protein)	phoR	2977	two-component sensor histidine kinase [PhoP]			
ytcB	3144	ABC transporter (ATP-binding protein)				cheV	1473	modulation of CheA activity in response to attractants (CheW and CheY similar domains)
ytcC	3143	ABC transporter (membrane protein)	resE	2416	two-component sensor histidine kinase [ResD]	cheW	1714	modulation of CheA activity in response to attractants
ythP	3071	ABC transporter (ATP-binding protein)	ybdK	222	two-component sensor histidine kinase [YbdJ]			
ytiC	3132	anion transport ABC transporter (ATP-binding protein)	ybcA	266	two-component sensor histidine kinase [YcbB]	flgB	1691	flagellar basal-body rod protein
			ybcM	279	two-component sensor histidine kinase [YcbL]	flgC	1691	flagellar basal-body rod protein
ytiD	3133	ABC transporter (permease)	ybcM	295	two-component sensor histidine kinase [YcbH]			
ytiP	3065	ABC transporter (permease)						

<i>flgA</i>	1700	flagellar hook protein	<i>coiX</i>	1251	spore coat protein (insoluble fraction)			SASP)
<i>flgB</i>	3639	flagellar hook-associated protein 1 (HAP1)	<i>coiY</i>	1250	spore coat protein (insoluble fraction)	<i>sspE</i>	937	small acid-soluble spore protein (major γ -type SASP)
<i>flgC</i>	3637	flagellar hook-associated protein 3 (HAP3)	<i>coiZ</i>	1249	spore coat protein (insoluble fraction)			
<i>flgM</i>	3640	flagellin synthesis regulatory protein (anti-sigma factor [σ^H])	<i>csqA</i>	228	sporulation-specific SASP protein	<i>sspF</i>	53	small acid-soluble spore protein (minor α/β -type SASP)
<i>flhA</i>	1707	flagella-associated protein	<i>jaq</i>	4213	SpolII-associated protein			
<i>flhB</i>	1706	flagella-associated protein	<i>kapB</i>	3230	activator of KinB in the initiation of sporulation	<i>usd</i>	3748	required for translation of <i>spoIIID</i>
<i>flhF</i>	1709	flagella-associated protein	<i>kapD</i>	3232	inhibitor of the KinA pathway to sporulation	<i>ynkT</i>	1495	sporulation protein σ^E -controlled
<i>flhO</i>	3746	flagellar basal-body rod protein	<i>kbaA</i>	159	activation of the KinB signaling pathway to sporulation	<i>ynkU</i>	1449	spore cortex membrane protein
<i>flhP</i>	3745	flagellar hook-basal body protein	<i>obg</i>	2853	GTP-binding protein involved in initiation of sporulation (SpoOA activation)	<i>ynzH</i>	1901	spore coat protein
<i>flhD</i>	3633	flagellar hook-associated protein 2 (HAP2)				<i>yobW</i>	2083	membrane protein σ^E -controlled
<i>flhE</i>	1652	flagellar hook-basal body protein	<i>phrA</i>	1316	phosphatase (RapA) inhibitor (imported by Opp)	<i>yagT</i>	2568	γ -glutamyl-L-di amino acid endopeptidase I
<i>flhG</i>	1652	flagellar basal-body M-ring protein	<i>phrC</i>	430	phosphatase (RapC) regulator / competence and sporulation stimulating factor (CSF)	<i>yagU</i>	2430	lipoprotein SpoIII-like
<i>flhH</i>	1694	flagellar assembly protein				<i>yraE</i>	2754	spore coat protein
<i>flhI</i>	1695	flagellar assembly protein	<i>phrE</i>	2660	phosphatase (RapE) regulator	<i>yraF</i>	2752	spore coat protein
<i>flhJ</i>	1695	flagellar-specific ATP synthase	<i>phrF</i>	3846	phosphatase (RapF) regulator	<i>yraG</i>	2752	spore coat protein
<i>flhK</i>	1697	flagellar protein required for formation of basal body	<i>phrG</i>	4141	phosphatase (RapG) regulator	<i>yraB</i>	2845	spore coat protein
<i>flhL</i>	1698	flagellar hook-length control	<i>phrI</i>	548	phosphatase (RapI) regulator	<i>yraC</i>	2844	spore coat protein
<i>flhM</i>	1701	flagellar protein required for flagellar formation	<i>phrK</i>	2063	phosphatase (RapK) regulator	<i>yraD</i>	2843	spore coat protein
<i>flhN</i>	1701	flagellar motor switch protein	<i>rapA</i>	1315	response regulator aspartate phosphatase [SpoOF-P]	<i>ytaP</i>	3161	spore coat protein
<i>flhP</i>	1704	flagellar protein required for flagellar formation	<i>rapB</i>	3771	response regulator aspartate phosphatase [SpoOF-P]	<i>ytpT</i>	3074	spore cortex protein
<i>flhQ</i>	1705	flagellar protein required for flagellar formation				<i>yyaA</i>	3051	DNA translocase stage III sporulation protein
<i>flhR</i>	1705	flagellar protein required for flagellar formation	<i>rapC</i>	428	response regulator aspartate phosphatase		4208	DNA-binding protein SpoII-like
<i>flhS</i>	1702	flagellar motor switch protein	<i>rapD</i>	3743	response regulator aspartate phosphatase	I.9	GERMINATION	23
<i>flhT</i>	1702	flagellar motor switch protein	<i>rapE</i>	2658	response regulator aspartate phosphatase	<i>gerAA</i>	3330	germination response to L-alanine
<i>flhZ</i>	1704	flagellar protein required for flagellar formation	<i>rapF</i>	3845	response regulator aspartate phosphatase	<i>gerAB</i>	3331	germination response to L-alanine
<i>hag</i>	3635	flagellin protein	<i>rapG</i>	4139	response regulator aspartate phosphatase	<i>gerAC</i>	3332	germination response to L-alanine
<i>mcpA</i>	3207	methyl-accepting chemotaxis protein (glucose and α -methyl-glucoside)	<i>rapH</i>	750	response regulator aspartate phosphatase	<i>gerBA</i>	3688	germination response to the combination of glucose, fructose, L-asparagine, and KCl
<i>mcpB</i>	3212	methyl-accepting chemotaxis protein (asparagine, glutamine and histidine)	<i>rapI</i>	547	response regulator aspartate phosphatase	<i>gerBB</i>	3689	germination response to the combination of glucose, fructose, L-asparagine, and KCl
<i>mcpC</i>	1463	methyl-accepting chemotaxis protein (cysteine, proline, threonine, glycine, serine, lysine, valine and arginine)	<i>rapK</i>	304	response regulator aspartate phosphatase	<i>gerBC</i>	3690	germination response to the combination of glucose, fructose, L-asparagine, and KCl
			<i>sinI</i>	2061	antagonist of SinR	<i>gerCB</i>	2384	heptaprenyl diphosphate synthase component I (menaquinone biosynthesis)
			<i>soj</i>	4206	centromere-like function involved in forespore chromosome partitioning / inhibition of SpoOA activation	<i>gerCC</i>	2382	heptaprenyl diphosphate synthase component II (menaquinone biosynthesis)
<i>motA</i>	1435	motility protein (flagellar motor rotation)	<i>spB</i>	1461	spore maturation protein (spore core dehydration)	<i>gerD</i>	159	germination response to L-alanine and to the combination of glucose, fructose, L-asparagine, and KCl
<i>motB</i>	1434	motility protein (flagellar motor rotation)	<i>spmA</i>	2423	spore maturation protein (spore core dehydration)	<i>gerKA</i>	420	germination response to the combination of glucose, fructose, L-asparagine, and KCl
<i>tipA</i>	3209	methyl-accepting chemotaxis protein	<i>spmB</i>	2422	spore maturation protein (spore core dehydration)	<i>gerKB</i>	423	germination response to the combination of glucose, fructose, L-asparagine, and KCl
<i>tipB</i>	3205	methyl-accepting chemotaxis protein				<i>gerKC</i>	421	germination response to the combination of glucose, fructose, L-asparagine, and KCl
<i>tipC</i>	374	methyl-accepting chemotaxis protein	<i>spo0B</i>	2854	sporulation initiation phosphoprotein (part of phosphorelay: Spo0F-P \rightarrow Spo0B-P \rightarrow Spo0A-P)	<i>gerM</i>	2902	germination response to L-alanine and to the combination of glucose, fructose, L-asparagine, and KCl
<i>yfmS</i>	808	methyl-accepting chemotaxis protein	<i>spo0E</i>	1430	negative sporulation regulatory phosphatase [SpoOA-P]	<i>gpr</i>	2635	spore protease (degradation of SASPs)
<i>yhlV</i>	1113	methyl-accepting chemotaxis protein	<i>spo0I</i>	4206	chromosome positioning near the pole and transport through the polar septum / antagonist of <i>Soj</i> anti-anti-sigma factor [SpoIAB]	<i>sleB</i>	2399	spore cortex-lytic enzyme
<i>yqgH</i>	1679	flagellar biosynthetic protein	<i>spollAA</i>	2444	anti-anti-sigma factor [SpoIAC]	<i>ykQ</i>	850	spore germination response
<i>yqgY</i>	1699	flagellar hook assembly protein	<i>spollAB</i>	2444	anti-anti-sigma factor [σ^H (SpoIAC)] and serine kinase [SpoIIA]	<i>ykR</i>	848	spore germination protein
<i>yxhH</i>	1710	flagellar biosynthesis switch protein	<i>spollB</i>	2864	endospore development (oligosporogenous mutation)	<i>ykT</i>	847	spore germination protein
<i>yxoH</i>	2030	methyl-accepting chemotaxis protein	<i>spollD</i>	3777	required for complete dissolution of the asymmetric septum	<i>ykvT</i>	1448	spore cortex-lytic enzyme
<i>yxD</i>	3043	flagellar motor apparatus	<i>spollE</i>	71	serine phosphatase [SpoIIA-P] (σ^E activation) / asymmetric septum formation	<i>yndD</i>	1907	spore germination protein
<i>yxE</i>	3042	motility protein	<i>spollGA</i>	1603	protease (processing of pro- σ^E to active σ^E)	<i>yndF</i>	1908	spore germination protein
<i>yxAQ</i>	3457	transmembrane receptor taxis protein	<i>spollIA</i>	2537	mutants block sporulation after engulfment		1909	spore germination protein
<i>yyvC</i>	3634	flagellar protein	<i>spollAB</i>	2536	mutants block sporulation after engulfment			
<i>yyvF</i>	3640	flagellar protein	<i>spollAD</i>	2535	mutants block sporulation after engulfment			
<i>yyvG</i>	3639	flagellar protein	<i>spollAE</i>	2535	mutants block sporulation after engulfment			
<i>yobE</i>	3609	flagellin	<i>spollAF</i>	2534	mutants block sporulation after engulfment			
			<i>spollAG</i>	2533	mutants block sporulation after engulfment			
			<i>spollAH</i>	2532	mutants block sporulation after engulfment			
			<i>spollIE</i>	1752	DNA translocase required for chromosome partitioning through the septum into the forespore			
			<i>spollII</i>	4214	essential for σ^E activity at stage III			
			<i>spollM</i>	2450	required for dissolution of the septal cell wall			
			<i>spollP</i>	2634	required for dissolution of the septal cell wall			
			<i>spollQ</i>	3760	required for completion of engulfment			
			<i>spollR</i>	3794	required for processing of pro- σ^E			
			<i>spollS</i>	1349	lethal when synthesized during vegetative growth in the absence of SpoIIB			
			<i>spollSB</i>	1348	disruption blocks sporulation after septum formation			
			<i>spolVA</i>	2387	required for proper spore cortex formation and coat assembly			
			<i>spolVB</i>	2520	intercompartmental signalling of pro- σ^E processing/activation in the mother-cell			
			<i>spolVCA</i>	2654	site-specific DNA recombinase required for creating the <i>sigK</i> gene (excision of the <i>skin</i> element)			
			<i>spolVFA</i>	2857	inhibitor of SpoIVFB			
			<i>spolVFB</i>	2856	protease (processing of pro- σ^E to active σ^E)			
			<i>spolVAA</i>	2443	mutants lead to the production of immature spores			
			<i>spolVAB</i>	2442	mutants lead to the production of immature spores			
			<i>spolVAC</i>	2441	mutants lead to the production of immature spores			
			<i>spolVAD</i>	2441	mutants lead to the production of immature spores			
			<i>spolVAE</i>	2440	mutants lead to the production of immature spores			
			<i>spolVAF</i>	2439	mutants lead to the production of immature spores			
			<i>spolVB</i>	2829	involved in spore cortex synthesis			
			<i>spolVC</i>	60	thermosensitive mutant blocks spore coat formation			
			<i>spolVE</i>	1590	required for spore cortex synthesis			
			<i>spolVFA</i>	1744	dipicolinate synthase subunit A			
			<i>spolVFB</i>	1745	dipicolinate synthase subunit B			
			<i>spolVG</i>	56	required for spore cortex synthesis			
			<i>spolVID</i>	2872	required for assembly of the spore coat			
			<i>spolVK</i>	1873	disruption leads to the production of immature spores			
			<i>spolVM</i>	1655	required for normal spore cortex and coat synthesis			
			<i>spolVR</i>	1015	involved in spore cortex synthesis			
			<i>spolVS</i>	1769	required for dehydration of the spore core and assembly of the coat			
			<i>spaA</i>	3892	spore coat polysaccharide synthesis			
			<i>spaB</i>	3891	spore coat polysaccharide synthesis			
			<i>spaC</i>	3890	spore coat polysaccharide synthesis			
			<i>spaD</i>	3889	spore coat polysaccharide synthesis			
			<i>spaE</i>	3888	spore coat polysaccharide synthesis			
			<i>spaF</i>	3887	spore coat polysaccharide synthesis			
			<i>spaG</i>	3886	spore coat polysaccharide synthesis			
			<i>spaH</i>	3885	spore coat polysaccharide synthesis			
			<i>spaI</i>	3884	spore coat polysaccharide synthesis			
			<i>spaK</i>	3883	spore coat polysaccharide synthesis			
			<i>spaL</i>	3025	small acid-soluble spore protein (major α -type SASP)			
			<i>sspA</i>	1050	small acid-soluble spore protein (major β -type SASP)			
			<i>sspC</i>	2155	small acid-soluble spore protein (minor α/β -type SASP)			
			<i>sspD</i>	1413	small acid-soluble spore protein (minor α/β -type SASP)			

<i>flhA</i>	1707	flagella-associated protein	<i>jaq</i>	4213	SpolII-associated protein			
<i>flhB</i>	1706	flagella-associated protein	<i>kapB</i>	3230	activator of KinB in the initiation of sporulation	<i>usd</i>	3748	required for translation of <i>spoIIID</i>
<i>flhF</i>	1709	flagella-associated protein	<i>kapD</i>	3232	inhibitor of the KinA pathway to sporulation	<i>ynkT</i>	1495	sporulation protein σ^E -controlled
<i>flhO</i>	3746	flagellar basal-body rod protein	<i>kbaA</i>	159	activation of the KinB signaling pathway to sporulation	<i>ynkU</i>	1449	spore cortex membrane protein
<i>flhP</i>	3745	flagellar hook-basal body protein	<i>obg</i>	2853	GTP-binding protein involved in initiation of sporulation (SpoOA activation)	<i>ynzH</i>	1901	spore coat protein
<i>flhD</i>	3633	flagellar hook-associated protein 2 (HAP2)				<i>yobW</i>	2083	membrane protein σ^E -controlled
<i>flhE</i>	1652	flagellar hook-basal body protein	<i>phrA</i>	1316	phosphatase (RapA) inhibitor (imported by Opp)	<i>yagT</i>	2568	γ -glutamyl-L-di amino acid endopeptidase I
<i>flhG</i>	1652	flagellar basal-body M-ring protein	<i>phrC</i>	430	phosphatase (RapC) regulator / competence and sporulation stimulating factor (CSF)	<i>yagU</i>	2430	lipoprotein SpoIII-like
<i>flhH</i>	1694	flagellar assembly protein				<i>yraE</i>	2754	spore coat protein
<i>flhI</i>	1695	flagellar assembly protein	<i>phrE</i>	2660	phosphatase (RapE) regulator	<i>yraF</i>	2752	spore coat protein
<i>flhJ</i>	1695	flagellar-specific ATP synthase	<i>phrF</i>	3846	phosphatase (RapF) regulator	<i>yraG</i>	2752	spore coat protein
<i>flhK</i>	1697	flagellar protein required for formation of basal body	<i>phrG</i>	4141	phosphatase (RapG) regulator	<i>yraB</i>	2845	spore coat protein
<i>flhL</i>	1698	flagellar hook-length control	<i>phrI</i>	548	phosphatase (RapI) regulator	<i>yraC</i>	2844	spore coat protein
<i>flhM</i>	1701	flagellar protein required for flagellar formation	<i>phrK</i>	2063	phosphatase (RapK) regulator	<i>yraD</i>	2843	spore coat protein
<i>flhN</i>	1701	flagellar motor switch protein	<i>rapA</i>	1315	response regulator aspartate phosphatase [SpoOF-P]	<i>ytaP</i>	3161	spore coat protein
<i>flhP</i>	1704	flagellar protein required for flagellar formation	<i>rapB</i>	3771	response regulator aspartate phosphatase [SpoOF-P]	<i>ytpT</i>	3074	spore cortex protein
<i>flhQ</i>	1705	flagellar protein required for flagellar formation				<i>yyaA</i>	3051	DNA translocase stage III sporulation protein
<i>flhR</i>	1705	flagellar protein required for flagellar formation	<i>rapC</i>	428	response regulator aspartate phosphatase		4208	DNA-binding protein SpoII-like
<i>flhS</i>	1702	flagellar motor switch protein	<i>rapD</i>	3743	response regulator aspartate phosphatase	I.9	GERMINATION	23
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<i>flhZ</i>	1704	flagellar protein required for flagellar formation	<i>rapF</i>	3845	response regulator aspartate phosphatase	<i>gerAB</i>	3331	germination response to L-alanine
<i>hag</i>	3635	flagellin protein	<i>rapG</i>	4139	response regulator aspartate phosphatase	<i>gerAC</i>	3332	germination response to L-alanine
<i>mcpA</i>	3207	methyl-accepting chemotaxis protein (glucose and α -methyl-glucoside)	<i>rapH</i>	750	response regulator aspartate phosphatase	<i>gerBA</i>	3688	germination response to the combination of glucose, fructose, L-asparagine, and KCl
<i>mcpB</i>	3212	methyl-accepting chemotaxis protein (asparagine, glutamine and histidine)	<i>rapI</i>	547	response regulator aspartate phosphatase	<i>gerBB</i>	3689	germination response to the combination of glucose, fructose, L-asparagine, and KCl
<i>mcpC</i>	1463	methyl-accepting chemotaxis protein (cysteine, proline, threonine, glycine, serine, lysine, valine and arginine)	<i>rapK</i>	304	response regulator aspartate phosphatase	<i>gerBC</i>	3690	germination response to the combination of glucose, fructose, L-asparagine, and KCl
			<i>sinI</i>	2061	antagonist of SinR	<i>gerCB</i>	2384	heptaprenyl diphosphate synthase component I (menaquinone biosynthesis)
			<i>soj</i>	4206	centromere-like function involved in forespore chromosome partitioning / inhibition of SpoOA activation	<i>gerCC</i>	2382	heptaprenyl diphosphate synthase component II (menaquinone biosynthesis)
<i>motA</i>	1435	motility protein (flagellar motor rotation)	<i>spB</i>	1461	spore maturation protein (spore core dehydration)	<i>gerD</i>	159	germination response to L-alanine and to the combination of glucose, fructose, L-asparagine, and KCl
<i>motB</i>	1434	motility protein (flagellar motor rotation)	<i>spmA</i>	2423	spore maturation protein (spore core dehydration)	<i>gerKA</i>	420	germination response to the combination of glucose, fructose, L-asparagine, and KCl
<i>tipA</i>	3209	methyl-accepting chemotaxis protein	<i>spmB</i>	2422	spore maturation protein (spore core dehydration)	<i>gerKB</i>	423	germination response to the combination of glucose, fructose, L-asparagine, and KCl
<i>tipB</i>	3205	methyl-accepting chemotaxis protein				<i>gerKC</i>	421	germination response to the combination of glucose, fructose, L-asparagine, and KCl
<i>tipC</i>	374	methyl-accepting chemotaxis protein	<i>spo0B</i>	2854	sporulation initiation phosphoprotein (part of phosphorelay: Spo0F-P \rightarrow Spo0B-P \rightarrow Spo0A-P)	<i>gerM</i>	2902	germination response to L-alanine and to the combination of glucose, fructose, L-asparagine, and KCl
<i>yfmS</i>	808	methyl-accepting chemotaxis protein	<i>spo0E</i>	1430	negative sporulation regulatory phosphatase [SpoOA-P]	<i>gpr</i>	2635	spore protease (degradation of SASPs)
<i>yhlV</i>	1113	methyl-accepting chemotaxis protein	<i>spo0I</i>	4206	chromosome positioning near the pole and transport through the polar septum / antagonist of <i>Soj</i> anti-anti-sigma factor [SpoIAB]	<i>sleB</i>	2399	spore cortex-lytic enzyme
<i>yqgH</i>	1679	flagellar biosynthetic protein	<i>spollAA</i>	2444	anti-anti-sigma factor [SpoIAC]	<i>ykQ</i>	850	spore germination response
<i>yqgY</i>	1699	flagellar hook assembly protein	<i>spollAB</i>	2444	anti-anti-sigma factor [σ^H (SpoIAC)] and serine kinase [SpoIIA]	<i>ykR</i>	848	spore germination protein
<i>yxhH</i>	1710	flagellar biosynthesis switch protein	<i>spollB</i>	2864	endospore development (oligosporogenous mutation)	<i>ykT</i>	847	spore germination protein
<i>yxoH</i>	2030	methyl-accepting chemotaxis protein	<i>spollD</i>	3777	required for complete dissolution of the asymmetric septum	<i>ykvT</i>	1448	spore cortex-lytic enzyme
<i>yxD</i>	3043	flagellar motor apparatus	<i>spollE</i>	71	serine phosphatase [SpoIIA-P] (σ^E activation) / asymmetric septum formation	<i>yndD</i>	1907	spore germination protein
<i>yxE</i>	3042	motility protein	<i>spollGA</i>	1603	protease (processing of pro- σ^E to active σ^E)	<i>yndF</i>	1908	spore germination protein
<i>yxAQ</i>	3457	transmembrane receptor taxis protein	<i>spollIA</i>	2537	mutants			

glsH	4033	β-glucosidase (cellulose degradation)	yjmA	1300	glucuronate isomerase	mngD	2510	citrate synthase III
bgIS	4011	endo-1-3-1,4 glucanase (lichenan degradation)	yjmD	1304	sorbitol dehydrogenase	odhA	2111	2-oxoglutarate dehydrogenase (E1 subunit)
crn	3569	catabolite repression HPr-like protein	yjme	1305	o-mannanot hydrolase	odhB	2108	2-oxoglutarate dehydrogenase (dihydrolipoamide transsuccinylase, E2 subunit)
csrA	3635	carbon storage regulator	yjmf	1306	2-deoxy-o-glucanate 3-dehydrogenase	sdhA	2907	succinate dehydrogenase (flavoprotein subunit)
frbB	1508	fructose 1-phosphate kinase	yjml	1311	altronate hydrolase	sdhB	2905	succinate dehydrogenase (iron-sulphur protein)
galE	3990	UDP-glucose 4-epimerase (galactose metabolism)	yjmc	1356	dolichol phosphate mannose synthase	sdhC	2908	succinate dehydrogenase (cytochrome b ₅₅₈ subunit)
galK	3921	galactokinase (galactose metabolism)	yjkb	1366	chloromuconate cycloisomerase	sucC	1680	succinyl-CoA synthetase (β subunit)
galT	3919	galactanase 1-phosphate uridylyltransferase (galactose metabolism)	yjkc	1367	polyisugr degrading enzyme	sucD	1681	succinyl-CoA synthetase (α subunit)
gdh	445	glucose 1-dehydrogenase	yjkrV	1427	ribulose-bisphosphate carboxylase	yjmc	1303	malate dehydrogenase
glcK	2571	glucose kinase	yjkl	1537	myo-inositol-1 or 4-monophosphatase	yqkI	2452	malate dehydrogenase
glgA	3167	start (bacterial glycogen) synthase (glycogen biosynthesis)	yjku	1477	glucose 1-dehydrogenase	ytsI	2930	malate dehydrogenase
glgB	3171	1,4- glucan branching enzyme (glycogen biosynthesis)	yjko	1442	glucose 1-dehydrogenase	ywkA	3801	malate dehydrogenase
glgC	3169	glucose-1-phosphate adenylyltransferase (glycogen biosynthesis)	yjor	1653	ribulose-5-phosphate 3-epimerase			
glgD	3168	required for glycogen biosynthesis	yjY	1741	deacetylase			
glpD	1004	glycerol-3-phosphate dehydrogenase (glycerol utilization)	yjnf	1943	endo-xylanase	ald	3277	L-alanine dehydrogenase
glpK	1003	glycerol kinase (glycerol utilization)	yjng	1951	propionyl-CoA carboxylase	ampS	1516	aminopeptidase
gnrK	4113	6-phospho- glucosidase (arbutin fermentation)	yjoc	2023	xylulokinase	ansA	2456	L-asparaginase
gnrZ	4116	6-phosphogluconate dehydrogenase (gluconate utilization)	yjod	2024	phosphoglycerate dehydrogenase	ansB	2455	L-aspartase
gpsA	2389	NAD(P)H-dependent glycerol-3-phosphate dehydrogenase	yjoe	2025	formate dehydrogenase	aprE	1105	extracellular alkaline serine protease (subtilisin E)
gutB	667	sorbitol dehydrogenase	yjof	2031	4-hydroxyphenylacetate-3-hydroxylase	aprX	1862	intracellular alkaline serine protease
iolB	4082	myo-inositol catabolism	yjog	2007	alcohol dehydrogenase	argB	1197	N-acetylglutamate 5-phosphotransferase (arginine biosynthesis)
iolC	4081	myo-inositol catabolism	yjoi	2507	phosphoenolpyruvate mutase	argC	1195	N-acetylglutamate 5-semialdehyde dehydrogenase (arginine biosynthesis)
iolD	4080	myo-inositol catabolism	yjod	2488	apocit-4-CoA carboxylase	argD	1198	N-acetylornithine aminotransferase (arginine biosynthesis)
iolE	4078	myo-inositol catabolism	yjrh	2780	formate dehydrogenase	argE	2142	acetylornithine deacetylase (arginine biosynthesis)
iolG	4076	myo-inositol 2-dehydrogenase (inositol catabolism)	yjrh	2778	methyltransferase	argF	1203	ornithine carbamoyltransferase (arginine biosynthesis)
iolH	4075	myo-inositol catabolism	yjri	2768	cyclodextrin metabolism	argG	3013	argininosuccinate synthase (arginine biosynthesis)
iolI	4074	myo-inositol catabolism	yjrp	2742	sugar-phosphate dehydrogenase	argH	3012	argininosuccinate lyase (arginine biosynthesis)
iolJ	4084	myo-inositol catabolism	yjrc	2950	endo-1,4- glucanase	argI	1196	ornithine acetyltransferase / amino-acid acetyltransferase (arginine biosynthesis)
kdgA	2323	deoxyphosphogluconate aldolase (pectin utilization)	yjse	2932	glycolate oxidase subunit	aroA	3046	3-deoxy-α-arabino-heptulosonate 7-phosphate synthase / chorismate mutase-isozyme 3 (shikimate pathway)
kdgK	2324	2-keto-3-deoxygluconate kinase (pectin utilization)	yjsh	2934	glycolate oxidase subunit	aroB	2378	3-dehydroquininate synthase (shikimate pathway)
kduD	2326	2-keto-3-deoxygluconate oxidoreductase (pectin utilization)	yjst	2969	plant metabolite dehydrogenase	aroC	2413	3-dehydroquininate dehydratase (shikimate pathway)
kduI	2325	5-keto-4-deoxyuronate isomerase (pectin utilization)	yjtl	2989	acetyl-CoA carboxylase	aroD	2645	shikimate 5-dehydrogenase (shikimate pathway)
lacA	3504	β-galactosidase	yjuf	3227	dihydrolipoamide E-acyltransferase	aroE	2368	5-enolpyruvylshikimate-3-phosphate synthase (shikimate pathway)
lclE	329	γ-lactate dehydrogenase	yjvg	3224	NADH-dependent butanol dehydrogenase	aroF	2380	chorismate synthase (shikimate pathway)
lclH	3959	6-phospho- glucosidase	yjyh	3222	NADH-dependent butanol dehydrogenase	aroH	2377	chorismate mutase (isozymes 1 and 2) (aromatic amino acids biosynthesis)
lipD	782	hydrolytic enzyme	yjyk	3224	NADH-dependent butanol dehydrogenase	arol	340	shikimate kinase (shikimate pathway)
melA	3100	α-D-galactoside galactohydrolase	yjyl	3215	exo-1,4-glucosidase	asd	1745	aspartate-semialdehyde dehydrogenase
mtiD	4051	mannitol-1-phosphate dehydrogenase	yjzm	3200	rhannulokinase	ask	2910	aspartokinase II attenuator
nagA	3594	N-acetylglucosamine-6-phosphate deacetylase (N-acetyl glucosamine utilization)	yjzn	3198	L-rhamnose isomerase	asb	3127	asparagine synthetase
nagB	3596	N-acetylglucosamine-6-phosphate isomerase (N-acetyl glucosamine utilization)	yjzr	3382	retinol dehydrogenase	asnH	4098	asparagine synthetase
narQ	3773	required for formate dehydrogenase activity	yjzr	3318	retinol dehydrogenase	aspB	2348	aspartate aminotransferase
pelB	2058	pectate lyase	yjzr	3318	retinol dehydrogenase	bcsA	2317	naringenin-chalcone synthase (phenylalanine metabolism)
pelD	2063	mannose-6-phosphate isomerase	yjzr	3318	retinol dehydrogenase	bfmBAA	2499	branched-chain α-keto acid dehydrogenase E1 (2-oxoisovalerate dehydrogenase α subunit)
pps	2035	phosphoenolpyruvate synthase	yjvV	3493	hydrolase	bfmBAB	2498	branched-chain α-keto acid dehydrogenase E1 (2-oxoisovalerate dehydrogenase β subunit)
pta	3865	phosphotransacetylase	yjvN	3427	plant-metabolite dehydrogenase	bfmBB	2497	branched-chain α-keto acid dehydrogenase E2 subunit (lipamide acyltransferase)
ptsH	1459	histidine-containing phosphocarrier protein of the phosphotransferase system (PTS) (HPr protein)	yjwC	3615	pyruvate, water dikinase	bItD	2718	spermine / spermidine acetyltransferase
rbtK	3701	ribokinase (ribose metabolism)	yjwF	3592	phosphoglycolate phosphatase	bpr	1599	bacillopeptidase F
sacA	3902	sucrase-6-phosphate hydrolase	yjvP	3591	O-acetyltransferase	cad	1535	lysine decarboxylase
sacB	3535	levansucrase	yjvH	3590	pectate lyase	carA	1199	carbamoyl-phosphate transferase-arginine (sub-unit A) (arginine biosynthesis)
sacC	2759	levanase	yjvH	3664	UDP-N-acetylglucosamine 2-epimerase	carB	1200	carbamoyl-phosphate transferase-arginine (sub-unit B) (arginine biosynthesis)
sacX	3941	negative regulatory protein of SacY	yjvD	3895	aldehyde dehydrogenase	ctpA	2133	carboxy-terminal processing protease
treA	891	trehalose-6-phosphate hydrolase	yjvD	3872	glucose 1-dehydrogenase	cysE	113	serine acetyltransferase (cysteine biosynthesis)
xta	251	β-xylosidase / α-L-arabinosidase (xylan degradation)	yjvD	3850	glycerol-inducible protein	cysH	1630	phosphoadenosine phosphosulfate reductase (cysteine biosynthesis)
xyIA	1891	xyllose isomerase (xyllose metabolism)	yjvF	3805	glycerol-inducible protein	cysK	82	cysteine synthetase A (cysteine biosynthesis)
xyIB	1893	xylulose kinase (xyllose metabolism)	yjvF	3730	NDP-sugar dehydrogenase	dal	517	D-alanine racemase
xynA	2054	endo-1,4-xylanase (xylan degradation)	yjvD	4091	glucose 1-dehydrogenase	dapA	1748	dihydrodipicolinate synthase (diaminopimelate / lysine biosynthesis)
xynB	1888	xylan β-1,4-xylosidase (xylan degradation)	yjvA	4040	endo-1,5-L-arabinosidase	dapG	2359	dihydrodipicolinate reductase (diaminopimelate / lysine biosynthesis)
xynD	1945	endo-1,4-xylanase (xylan degradation)	yjvA	4000	gluconate 5-dehydrogenase	def	1646	polypeptide deformylase
ybaN	161	polysaccharide deacetylase	yjvA	4107	glucose 1-dehydrogenase	epo	3939	minor extracellular serine protease
ybbD	188	β-hexosaminidase	yjvA	4202	formate dehydrogenase	gImS	200	L-glutamine-D-fructose-6-phosphate amidotransferase
ybcM	213	glucosamine-fructose-6-phosphate aminotransferase	yjvA	4196	galactoside acetyltransferase	glnA	1878	glutamine synthetase
ybtT	258	glucosamine-6-phosphate isomerase	yjvA	4136	formaldehyde dehydrogenase	gltA	2014	glutamate synthase (large subunit) (glutamate biosynthesis)
yctB	268	5-dehydro-D-deoxyglucarate dehydratase	iln2			gltB	2009	glutamate synthase (small subunit) (glutamate biosynthesis)
yctC	269	acetylaldehyde dehydrogenase	iln3			gltG	3789	serine hydroxymethyltransferase (glycine / serine / threonine metabolism)
yctF	272	glutamate dehydratase	iln3			hisA	3584	phosphoribosylformimino-5-aminoimidazole carboxamide ribotide isomerase (histidine biosynthesis)
ydcF	305	glucose 1-dehydrogenase	iln3			hisB	3585	imidazoleglycerol-phosphate dehydratase (histidine biosynthesis)
ydcG	306	oligo-1,6-glucosidase	iln3			hisC	2371	glutathionyl-phosphate aminotransferase (histidine biosynthesis) / tyrosine and phenylalanine aminotransferase
ydcS	352	aromatic hydrocarbon catabolism	iln3			hisD	3587	histidinol dehydrogenase (histidine biosynthesis)
ydcK	370	β-glucosidase	iln3			hisF	3583	HisF cyclase-like protein (synthesis of D-erythroimidazole glycerol phosphate)
ydcG	375	D-arabino-3-hexulose 6-phosphate formaldehyde lyase	iln3			hisG	3587	ATP phosphoribosyltransferase (histidine biosynthesis)
ydsN	466	aryl-alcohol dehydrogenase	iln3			hisH	3585	amidotransferase (histidine biosynthesis)
ydaD	471	alcohol dehydrogenase	iln3			hisI	3583	phosphoribosyl-AMP cyclohydrolase / phosphoribosyl-ATP pyrophosphohydrolase (histidine biosynthesis)
ydaF	473	acetyltransferase	iln3			hom	3315	homoserine dehydrogenase (threonine / methionine biosynthesis)
ydaM	482	cellulose synthase	iln3			hutG	4045	formingonlutamate hydrolase (histidine utilization)
ydaP	488	reticuline oxidase	iln3			hutH	4041	histidase (histidine utilization)
ydhR	628	β-glucosidase	iln3			hutI	4044	imidazole-5-propionate hydrolase (histidine utilization)
ydhR	631	fructokinase	iln3			hutU	4042	urocanase (histidine utilization)
ydhS	632	mannose-6-phosphate isomerase	iln3			ilvA	2896	threonine dehydratase (isoleucine biosynthesis)
ydhT	632	mannan endo-1,4-mannosidase	iln3			ilvB		acetolactate synthase (large subunit) (valine / isoleucine biosynthesis)
ydhE	670	fructokinase	iln3			ilvC	2894	keto-acid reductoisomerase (valine / isoleucine biosynthesis)
ydhJ	679	L-ditol 2-dehydrogenase	iln3			ilvD	2302	dihydroxy-acid dehydratase (valine / isoleucine biosynthesis)
ydhP	682	arylesterase	iln3			ilvN	2894	acetolactate synthase (small subunit)
yecA	688	methanethiol dehydrogenase regulation	iln3					
yeyY	774	rhannogalacturonan acetyltransferase	iln3					
yeyZ	774	β-galactosidase	iln3					
yfhM	829	epoxide hydrolase	iln3					
yfhR	937	glucose 1-dehydrogenase	iln3					
yfS	969	polysaccharide deacetylase	iln3					
yfmT	807	benzaldehyde dehydrogenase	iln3					
yfhH	798	glucose 1-phosphate cytidylyltransferase	iln3					
ygaK	958	reticuline oxidase	iln3					
yhcW	997	phosphoglycolate phosphatase	iln3					
yhdF	1022	glucose 1-dehydrogenase	iln3					
yhdN	1030	aldo / keto reductase	iln3					
yheN	1041	endo-1,4-xylanase	iln3					
yhiE	1095	glucanase	iln3					
yhxH	1006	phosphomannomutase	iln3					
yhxC	1115	alcohol dehydrogenase	iln3					
yhbD	1110	alcohol dehydrogenase	iln3					
yisS	1164	myo-inositol 2-dehydrogenase	iln3					
yitS	1175	mandelate racemase	iln3					
yitY	1192	L-gulonolactone oxydase	iln3					
yidE	1274	mannose-6-phosphate isomerase	iln3					
yieA	1281	endo-1,4-xylanase	iln3					
yigC	1285	formate dehydrogenase	iln3					
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ipIA	4083	(valine/isoleucine biosynthesis)	yqiE	2486	tripeptidase	xpt	2319	xanthine phosphoribosyltransferase (purine biosynthesis)
ipo	1189	methylmalonate-semialdehyde dehydrogenase (valine metabolism)	yqiN	2475	amino acid degradation	yafF	23	deoxyphenyl kinase subunit
ispA	1386	intracellular proteinase inhibitor	yqiR	2472	pyrroline-5-carboxylate reductase	yaaG	24	deoxyphenyl kinase subunit
kbl	1771	2-amino-3-ketobutyrate CoA ligase	yriE	2470	D-serine dehydratase	yabR	70	polyribonucleotide nucleotidyltransferase
leuA	2893	2-isopropylmalate synthase (leucine biosynthesis)	yriH	2839	opine catabolism	yerA	713	adenine deaminase
leuB	2891	3-isopropylmalate dehydrogenase (leucine biosynthesis)	yriH	2785	cystathionine γ -synthase	yriK	859	2',3'-cyclic-nucleotide 2'-phosphodiesterase
leuC	2890	3-isopropylmalate dehydratase (large subunit) (leucine biosynthesis)	yriP	2768	dihydrodipicolinate reductase	yriM	1069	GMP-binding factor
leuD	2889	3-isopropylmalate dehydratase (small subunit) (leucine biosynthesis)	yriP	2738	glutamate racemase	yriR	991	5'-nucleotidase
lysA	2437	3-deoxyisopimelate decarboxylase (lysine biosynthesis)	yriN	2794	protease	yirY	1144	DNA exonuclease
lysC	2910	aspartokinase II (α and β subunits) (diaminopimelate/lysine biosynthesis)	yriO	2793	protease	yjdB	1236	GTP pyrophosphokinase
metB	2305	homoserine O-succinyltransferase (methionine biosynthesis)	yriE	2897	acetyltransferase	yjDP	1240	diadenosine tetraphosphatase
metC	1385	cobalamin-independent methionine synthase (methionine biosynthesis)	yriD	3146	N-acetylaminic acid racemase	yjKE	1377	formyltetrahydrofolate deformylase
metK	3128	S-adenosylmethionine synthetase	yriC	3066	cysteine synthase	yjLD	1585	IMP dehydrogenase
npr	245	extracellular metalloprotease	yriB	3193	cyano dioxygenase	yjLO	1641	guanylate kinase
nasB	362	intracellular nitrate reductase (electron transfer subunit)	yriH	3226	aspartate aminotransferase	yjMA	1868	ribonucleoprotein
nasC	360	assimilatory nitrate reductase (catalytic subunit)	yriG	3341	aspartate aminotransferase	yjNC	1895	micrococcal nuclease
nasD	358	assimilatory nitrite reductase (subunit)	yriH	3343	N-carbamyl-L-amino acid amidohydrolase	yjNF	1899	deoxyuridine 5'-triphosphate pyrophosphatase
nasE	355	assimilatory nitrite reductase (subunit)	yriH	3347	opine catabolism	yjNS	2165	ribonucleoside-diphosphate reductase (α subunit)
nprB	1186	extracellular neutral protease B	yriP	3351	opine catabolism	yjOS	2164	ribonucleoside-diphosphate reductase (α subunit)
nprE	1541	extracellular neutral metalloprotease	yriR	3353	opine catabolism	yjOS	2161	ribonucleoside-diphosphate reductase (β subunit)
nrgB	3757	nitrogen-regulated Pil-like protein	yriT	3354	methylglyoxalase	yjP	2159	uridylic acid 5'-triphosphate nucleotidohydrolase
patA	1472	aminotransferase	yriH	3366	glycine cleavage system protein H	yjPD	2395	ribosealdehyde protein S1 homologue
patB	3228	aminotransferase	yriM	3373	proline dehydrogenase	yjBE	2528	exodeoxyribonuclease VII (large subunit)
pepT	3994	peptidase T	yriX	3381	oligodeoxyribose epimerase	yjCI	2526	exodeoxyribonuclease VII (small subunit)
pheA	2851	phenylalanine dehydratase (phenylalanine biosynthesis)	yriL	3376	diaminopimelate epimerase	yjDF	2730	ribonuclease inhibitor
pheB	2852	chorismate mutase (phenylalanine biosynthesis)	yriL	3312	acylaminoacyl-peptidase	yjDT	2787	purine nucleoside phosphorylase
proA	1379	γ -glutamyl phosphate reductase (proline biosynthesis)	yriK	3452	carboxylesterase	yjMD	3202	GMP reductase
proB	1378	γ -glutamyl kinase (proline biosynthesis)	yriD	3516	serine O-acetyltransferase	yjNH	3328	allantoinase
proH	2017	involved in proline biosynthesis (salt-inducible)	yriB	3623	carboxy-terminal processing protease	yjNL	3332	uricase
proI	2016	glutamate 5-kinase (proline biosynthesis)	yriA	3956	branched-chain amino acid aminotransferase	yjR	3343	ribonuclease
racX	3533	amino acid racemase	yriD	3947	aminopeptidase	yjWC	3349	GTP-pyrophosphokinase
rocA	3879	proline 5-carboxylate dehydrogenase (arginine and ornithine utilization)	yriE	3881	glutamate dehydrogenase	ilA	77	METABOLISM OF LIPIDS
rocB	3878	involved in arginine and ornithine utilization	yriF	3868	aspartate aminotransferase	accB	2531	acetyl-CoA carboxylase (α subunit) (long-chain fatty acid biosynthesis)
rocD	4145	ornithine aminotransferase (arginine and ornithine utilization)	yriF	3849	sermidine synthase	accC	2531	acetyl-CoA carboxylase (biotin carboxyl carrier subunit) (long-chain fatty acid biosynthesis)
rocF	4142	arginase (arginine and ornithine utilization)	yriD	3848	glymatase	accD	3813	acetyl-CoA dehydrogenase
serA	2410	phosphoglycerate dehydrogenase (serine biosynthesis)	yriD	3720	γ -glutamyltransferase	accA	1665	acyl carrier protein (fatty acid biosynthesis)
serC	1076	phosphoserine aminotransferase (serine biosynthesis)	yriD	4057	aminoacylase	accA	1721	acyl carrier protein (fatty acid biosynthesis)
tdh	1770	threonine 3-dehydrogenase (threonine catabolism)	ilB	77	METABOLISM OF NUCLEOTIDES AND NUCLEIC ACIDS	accA	1721	acyl carrier protein (fatty acid biosynthesis)
thrB	3313	homoserine kinase (threonine biosynthesis)	adeC	1521	adenine deaminase	accA	1721	acyl carrier protein (fatty acid biosynthesis)
thrC	3314	threonine synthase (threonine biosynthesis)	adeC	1521	adenine deaminase	accA	1721	acyl carrier protein (fatty acid biosynthesis)
trpA	2372	tryptophan synthase (α subunit) (tryptophan biosynthesis)	adeC	1521	adenine deaminase	accA	1721	acyl carrier protein (fatty acid biosynthesis)
trpB	2373	tryptophan synthase (β subunit) (tryptophan biosynthesis)	ade					

<i>ywpB</i>	3743	hydroxymyristoyl-(acyl carrier protein) dehydrogenase	<i>yjbT</i>	1245	thiamin biosynthesis	<i>recR</i>	29	DNA repair and genetic recombination
<i>yxD</i>	4001	3-oxoadipate CoA-transferase	<i>yjbU</i>	1246	thiamin biosynthesis	<i>ruvA</i>	2636	Holliday junction DNA helicase
<i>yxE</i>	4001	3-oxoadipate CoA-transferase	<i>yjbV</i>	1246	thiamin biosynthesis	<i>ruvB</i>	2636	Holliday junction DNA helicase
			<i>ykpB</i>	1513	thiamin biosynthesis	<i>sbcD</i>	1143	endonuclease SbcD homologue
			<i>ykvK</i>	1440	6-pyruvoyl tetrahydrobiopterin synthase	<i>ylpB</i>	1659	ATP-dependent DNA helicase
II.5		METABOLISM OF COENZYMES AND PROSTHETIC GROUPS 99	<i>ykvL</i>	1440	coenzyme PQQ synthetase	<i>yocI</i>	2095	ATP-dependent DNA helicase
<i>bioA</i>	3094	adenosylmethionine-8-amino-7-oxononanoate aminotransferase (biotin biosynthesis)	<i>ykbQ</i>	1577	pyrimidine-thiamine biosynthesis	<i>yorkK</i>	2180	single-strand DNA-specific exonuclease
<i>bioB</i>	3091	biotin synthetase (biotin biosynthesis)	<i>ylnD</i>	1633	uroporphyrin-III C-methyltransferase	<i>yqhH</i>	2549	SNF2 helicase
<i>bioD</i>	3091	dethiobiotin synthetase (biotin biosynthesis)	<i>ylnF</i>	1635	uroporphyrin-III C-methyltransferase	<i>yrrC</i>	2808	conjugation transfer protein
<i>bioF</i>	3092	8-amino-7-oxononanoate synthase (biotin biosynthesis)	<i>yiol</i>	1642	pantothenate metabolism flavoprotein	<i>yrvE</i>	2825	single-strand DNA-specific exonuclease
<i>bioI</i>	3089	cytochrome P450-like enzyme (biotin biosynthesis)	<i>yngH</i>	1954	biotin carboxylase	<i>yvwA</i>	3735	SNF2 helicase
<i>bioW</i>	3094	6-carboxyhexanoate-CoA ligase (biotin biosynthesis)	<i>yodC</i>	2127	nitroreductase			
<i>dtfA</i>	2296	dihydrofolate reductase (glycine/purine/DNA precursor synthesis, conversion of dUMP to dTMP)	<i>yqgV</i>	2574	5-formyltetrahydrofolate cyclo-ligase	III.4		DNA PACKAGING AND SEGREGATION 10
<i>dhaS</i>	2100	aldehyde dehydrogenase	<i>yqS</i>	2469	pantothenate kinase	<i>grlA</i>	1935	DNA gyrase-like protein (subunit A)
<i>dhaB</i>	3291	2,3-dihydro-2,3-dihydroxybenzoate dehydrogenase (2,3-dihydroxybenzoate biosynthesis)	<i>yrrL</i>	2795	folate metabolism	<i>grlB</i>	1933	DNA gyrase-like protein (subunit B)
<i>dhaB</i>	3288	isochorismatase (2,3-dihydroxybenzoate biosynthesis)	<i>yrrM</i>	2795	caffeoyl-CoA O-methyltransferase	<i>gyrA</i>	7	DNA gyrase (subunit A)
<i>dhaC</i>	3291	isochorismatase synthase (2,3-dihydroxybenzoate biosynthesis)	<i>yueD</i>	3265	sepiapterin reductase	<i>gyrB</i>	5	DNA gyrase (subunit B)
<i>dhaE</i>	3289	2,3-dihydroxybenzoate-AMP ligase (enterobactin synthetase component E) (2,3-dihydroxybenzoate biosynthesis)	<i>yueI</i>	3261	pyrazinamidase/nicotinamidase	<i>hbs</i>	2385	non-specific DNA-binding protein HBSu
			<i>yueK</i>	3260	nicotinate phosphoribosyltransferase	<i>smc</i>	1666	chromosome segregation SMC protein homologue
<i>dhaF</i>	3287	involved in 2,3-dihydroxybenzoate biosynthesis	<i>yulG</i>	3293	biotin metabolism			
<i>folA</i>	87	7,8-dihydro-6-hydroxymethylpterin pyrophosphokinase (dihydrofolate biosynthesis)	<i>yurB</i>	3335	4-hydroxybenzoyl-CoA reductase	<i>smf</i>	1682	DNA processing Sml protein homologue
<i>folC</i>	2866	folyl-polyglutamate synthetase (folate biosynthesis)	<i>yurC</i>	3338	4-hydroxybenzoyl-CoA reductase	<i>topA</i>	1683	DNA topoisomerase I
<i>folD</i>	2529	methylene tetrahydrofolate dehydrogenase / methenyltetrahydrofolate cyclohydrolase (purines and amino acids biosynthesis)	<i>yurD</i>	3338	4-hydroxybenzoyl-CoA reductase	<i>topB</i>	476	DNA topoisomerase III
			<i>yutB</i>	3320	lipic acid synthetase	<i>yonN</i>	2225	HU-related DNA-binding protein
<i>folK</i>	87	7,8-dihydro-6-hydroxymethylpterin pyrophosphokinase (dihydrofolate biosynthesis)	<i>ywaB</i>	3950	quinone biosynthesis			
<i>ggT</i>	2004	γ -glutamyltranspeptidase (glutathione metabolism)	<i>ywkE</i>	3796	protoporphyrinogen oxidase			
			<i>ywoC</i>	3755	isochorismatase	III.5		RNA SYNTHESIS 244
<i>gsaB</i>	943	glutamate-1-semialdehyde aminotransferase	II.6		METABOLISM OF PHOSPHATE 9	III.5.1		INITIATION 19
<i>hemA</i>	2878	glutamyl-tRNA reductase (porphyrin biosynthesis)	<i>phoA</i>	1018	alkaline phosphatase A	<i>sigA</i>	2601	RNA polymerase major sigma factor (σ^70)
<i>hemB</i>	2874	8-aminolevulinic acid dehydratase (porphyrin biosynthesis)	<i>phoB</i>	621	alkaline phosphatase III	<i>sigB</i>	522	RNA polymerase general stress sigma factor (σ^32)
<i>hemC</i>	2876	porphobilinogen deaminase (porphyrin biosynthesis)	<i>phoD</i>	284	phosphodiesterase/alkaline phosphatase	<i>sigD</i>	1716	RNA polymerase flagella, motility, chemotaxis and autolysis sigma factor (σ^22)
<i>hemD</i>	2875	uroporphyrinogen III cosynthase (porphyrin biosynthesis)	<i>phoH</i>	2615	phosphate starvation-induced protein	<i>sigE</i>	1604	RNA polymerase sporulation mother cell-specific (early) sigma factor (σ^54) (SpoIIIG)
<i>hemE</i>	1086	uroporphyrinogen III decarboxylase (porphyrin biosynthesis)	<i>xpaC</i>	36	hydrolysis of 5-bromo-4-chloroindolyl phosphate	<i>sigF</i>	2443	RNA polymerase sporulation forespore-specific (early) sigma factor (σ^70) (SpoIIAC)
<i>hemH</i>	1087	ferrochelatase (porphyrin biosynthesis)	<i>yblM</i>	248	alkaline phosphatase	<i>sigG</i>	1605	RNA polymerase sporulation forespore-specific (late) sigma factor (σ^70) (SpoIIIG)
<i>hemL</i>	2873	glutamate-1-semialdehyde 2,1-aminotransferase (porphyrin biosynthesis)	<i>ykoX</i>	1409	alkaline phosphatase	<i>sigH</i>	117	RNA polymerase vegetative and early stationary-phase sigma factor (σ^70) (SpoIIH)
<i>hemN</i>	2630	coproporphyrinogen III oxidase (porphyrin biosynthesis)	<i>yiaK</i>	1549	phosphate starvation inducible protein	<i>sigI</i>	3513	RNA polymerase sigma factor (σ^70)
<i>hemX</i>	2877	negative effector of the concentration of HemA	<i>yngC</i>	1947	alkaline phosphatase	<i>sigV</i>	2769	RNA polymerase ECF-type sigma factor (σ^54)
<i>hemY</i>	1088	protoporphyrinogen IX oxidase (porphyrin biosynthesis)	II.7		METABOLISM OF SULPHUR 8	<i>sigW</i>	195	RNA polymerase ECF-type sigma factor (σ^54)
<i>menB</i>	3149	dihydroxynaphthoic acid synthetase (menaquinone biosynthesis)	<i>yisZ</i>	1170	adenylsulfate kinase	<i>sigX</i>	2414	RNA polymerase ECF-type sigma factor (σ^54)
<i>menD</i>	3151	2-succinyl-6-hydroxy-2,4-cyclohexadiene-1-carboxylate synthase / 2-oxoglutarate decarboxylase (menaquinone biosynthesis)	<i>yitA</i>	1171	sulfate adenyltransferase	<i>sigY</i>	3970	RNA polymerase ECF-type sigma factor (σ^54)
<i>menE</i>	3148	O-succinylbenzoic acid-CoA ligase (menaquinone biosynthesis)	<i>yitB</i>	1172	phospho-adenylsulfate sulfotransferase	<i>sigZ</i>	2742	RNA polymerase ECF-type sigma factor (σ^54)
<i>menF</i>	3153	menaquinone-specific isochorismatase synthase (menaquinone biosynthesis)	<i>yitC</i>	1173	phospho-adenylsulfate sulfotransferase	<i>spolIIC</i>	2701	RNA polymerase sporulation mother cell-specific (late) sigma factor (σ^70) (C-terminal half)
<i>moaB</i>	3014	molybdopterin precursor biosynthesis	<i>yitD</i>	1173	phospho-adenylsulfate sulfotransferase	<i>spolIIC</i>	2652	RNA polymerase sporulation mother cell-specific (late) sigma factor (σ^70) (N-terminal half)
<i>moaD</i>	1499	molybdopterin converting factor (subunit 1)	<i>yitE</i>	1173	phospho-adenylsulfate sulfotransferase	<i>xpf</i>	1324	RNA polymerase PBX sigma factor-like
<i>moaE</i>	1498	molybdopterin converting factor (subunit 2)	<i>yitF</i>	1172	phospho-adenylsulfate sulfotransferase	<i>yhdM</i>	1030	RNA polymerase ECF-type sigma factor
<i>moaF</i>	1495	molybdopterin-guanine dinucleotide biosynthesis	<i>yitG</i>	1172	phospho-adenylsulfate sulfotransferase	<i>ykoZ</i>	1411	RNA polymerase sigma factor
<i>moaB</i>	1498	molybdopterin-guanine dinucleotide biosynthesis	<i>yitH</i>	1173	phospho-adenylsulfate sulfotransferase	<i>yiaC</i>	1543	RNA polymerase ECF-type sigma factor
<i>moaF</i>	1497	molybdopterin biosynthesis protein	<i>yitI</i>	1173	phospho-adenylsulfate sulfotransferase			
<i>moaB</i>	1496	molybdopterin biosynthesis protein	<i>yitJ</i>	1173	phospho-adenylsulfate sulfotransferase	III.2		REGULATION 213
<i>mtA</i>	2385	GTP cyclohydrolase I (tetrahydrofolate biosynthesis)	<i>yitK</i>	1173	phospho-adenylsulfate sulfotransferase	<i>abh</i>	1517	transcriptional regulator of transition state genes (AbrB-like)
<i>nadA</i>	2846	quinolinate synthetase (quinolinate biosynthesis)	<i>yitL</i>	1173	phospho-adenylsulfate sulfotransferase	<i>abrB</i>	45	transcriptional pleiotropic regulator of transition state genes (<i>aprE</i> , <i>comK</i> , <i>ftsAZ</i> , <i>hpr</i> , <i>motAB</i> , <i>npfE</i> , <i>pbpE</i> , <i>rbp</i> , <i>spo0H</i> , <i>spoVG</i> , <i>spoVE</i> , <i>tycA</i>)
<i>nadB</i>	2849	L-aspartate oxidase (quinolinate biosynthesis)	<i>yitM</i>	1173	phospho-adenylsulfate sulfotransferase	<i>acoR</i>	883	transcriptional activator of the acetoin dehydrogenase operon (<i>acoABCD</i>)
<i>nadC</i>	2847	nicotinate-nucleotide pyrophosphorylase (NAD/NADP biosynthesis)	<i>yitN</i>	1173	phospho-adenylsulfate sulfotransferase	<i>ahrC</i>	2522	transcriptional regulator of arginine metabolism expression (<i>roc</i> operon)
<i>nadE</i>	338	NH ₄ ⁺ -dependent NAD ⁺ synthetase (NAD biosynthesis)	<i>yitO</i>	1173	phospho-adenylsulfate sulfotransferase	<i>alsR</i>	3711	transcriptional regulator of the α -acetolactate operon (<i>alsSD</i>)
<i>narA</i>	3772	molybdopterin precursor biosynthesis	<i>yitP</i>	1173	phospho-adenylsulfate sulfotransferase	<i>ansR</i>	2456	transcriptional repressor of the <i>ansAB</i> operon (<i>Xre</i> family)
<i>narX</i>	355	uroporphyrin-III C-methyltransferase (porphyrin biosynthesis)	<i>yitQ</i>	1173	phospho-adenylsulfate sulfotransferase	<i>araR</i>	3485	transcriptional repressor of the arabinose operon (<i>araBADLMNPO</i>)
<i>nifS</i>	2849	required for NAD biosynthesis	<i>yitR</i>	1173	phospho-adenylsulfate sulfotransferase	<i>azlB</i>	2729	transcriptional repressor of the <i>azlBCD</i> operon
<i>pabA</i>	84	p-aminobenzoate synthase glutamine amidotransferase (subunit B) / anthranilate synthase (subunit II) (folate and tryptophan biosynthesis)	<i>yitS</i>	1173	phospho-adenylsulfate sulfotransferase	<i>birA</i>	2355	transcriptional repressor of the biotin operon (<i>bioWAFDBI</i>) / biotin acetyl-CoA-carboxylase synthetase
<i>pabB</i>	83	p-aminobenzoate synthase (subunit A) (folate biosynthesis)	<i>yitT</i>	1173	phospho-adenylsulfate sulfotransferase	<i>bltR</i>	2716	transcriptional regulator of the <i>bltD</i> operon
<i>pabC</i>	85	aminodeoxychorismate lyase (folate biosynthesis)	<i>yitU</i>	1173	phospho-adenylsulfate sulfotransferase	<i>bmrR</i>	2495	transcriptional activator of the <i>bmrUR</i> operon
<i>panB</i>	2354	ketopantoate hydroxymethyltransferase (pantothenate biosynthesis)	<i>yitV</i>	1173	phospho-adenylsulfate sulfotransferase	<i>ccpA</i>	3044	transcriptional regulator involved in carbon catabolite control
<i>panC</i>	2353	pantothenate synthetase (pantothenate biosynthesis)	<i>yitW</i>	1173	phospho-adenylsulfate sulfotransferase	<i>cheB</i>	1711	two-component response regulator-like [CheA] / methyl-accepting chemotaxis proteins-glutamate methyltransferase
<i>panD</i>	2352	aspartate 1-decarboxylase (pantothenate biosynthesis)	<i>yitX</i>	1173	phospho-adenylsulfate sulfotransferase	<i>cheY</i>	1703	two-component response regulator [CheA] involved in modulation of flagellar switch bias (chemotaxis)
<i>ribA</i>	2429	GTP cyclohydrolase II / 3,4-dihydroxy-2-butanone 4-phosphate synthase (riboflavin biosynthesis)	<i>yitY</i>	1173	phospho-adenylsulfate sulfotransferase	<i>citR</i>	1020	transcriptional repressor of the citrate synthase I gene (<i>citA</i>)
<i>ribB</i>	2429	riboflavin synthase (α subunit) (riboflavin biosynthesis)	<i>yitZ</i>	1173	phospho-adenylsulfate sulfotransferase	<i>citT</i>	832	two-component response regulator [CitS]
<i>ribC</i>	1737	riboflavin kinase / FAD synthase (riboflavin biosynthesis)	<i>yitA</i>	1173	phospho-adenylsulfate sulfotransferase	<i>codY</i>	1690	transcriptional pleiotropic repressor (expression of <i>srfA</i> , <i>comK</i> , <i>dpp</i> , <i>gabP</i> , <i>hut</i> , <i>ureABC</i>)
<i>ribG</i>	2431	riboflavin-specific deaminase (riboflavin biosynthesis)	<i>yitB</i>	1173	phospho-adenylsulfate sulfotransferase	<i>comA</i>	3253	two-component response regulator [ComP] of late competence genes / surfactin production
<i>ribH</i>	2428	riboflavin synthase (β subunit) (riboflavin biosynthesis)	<i>yitC</i>	1173	phospho-adenylsulfate sulfotransferase	<i>comK</i>	1117	competence transcription factor (CTF), final autoregulatory control switch prior to competence development
<i>ribT</i>	2427	reductase (riboflavin biosynthesis)	<i>yitD</i>	1173	phospho-adenylsulfate sulfotransferase	<i>comQ</i>	3256	transcriptional regulator of late competence operon (<i>comG</i>) and surfactin expression (<i>srfA</i>)
<i>sul</i>	86	dihydropterate synthase (dihydrofolate biosynthesis)	<i>yitE</i>	1173	phospho-adenylsulfate sulfotransferase	<i>ctsR</i>	101	transcriptional repressor of class III stress genes (<i>clpC</i> , <i>clpP</i>)
<i>thiA</i>	955	synthesis of the pyrimidine moiety of thiamin (thiamin biosynthesis)	<i>yitF</i>	1173	phospho-adenylsulfate sulfotransferase	<i>degA</i>	1163	transcriptional activator involved in the degradation of glutamine phosphoribosylpyrophosphate amidotransferase
<i>thiC</i>	3930	thiamine-phosphate pyrophosphorylase (thiamin biosynthesis)	<i>yitG</i>	1173	phospho-adenylsulfate sulfotransferase	<i>degU</i>	3644	two-component response regulator [DegS] involved in degradative enzyme and competence regulation (<i>sacB</i> , <i>degQ</i> , <i>comK</i>)
<i>thiD</i>	3930	phosphomethylpyrimidine kinase (thiamin biosynthesis)	<i>yitH</i>	1173	phospho-adenylsulfate sulfotransferase	<i>deoR</i>	4052	transcriptional repressor of the <i>draI</i> / <i>nupC</i> / <i>pdp</i> operon (deoxyribonucleoside)
<i>thiK</i>	3931	hydroxyethylthiazole kinase (thiamin biosynthesis)	<i>yitI</i>	1173	phospho-adenylsulfate sulfotransferase	<i>fnr</i>	3831	transcriptional regulator of anaerobic genes (<i>narK</i> , <i>narGHI</i>)
<i>yaal</i>	26	isochorismatase	<i>yitJ</i>	1173	phospho-adenylsulfate sulfotransferase	<i>fruR</i>	1507	transcriptional repressor of the fructose operon (<i>fruRBA</i>)
<i>ydiA</i>	640	thiamin-monophosphate kinase	<i>yitK</i>	1173	phospho-adenylsulfate sulfotransferase	<i>gerE</i>	2904	transcriptional regulator required for expression of late spore coat genes
<i>ydiG</i>	646	molybdopterin precursor biosynthesis	<i>yitL</i>	1173	phospho-adenylsulfate sulfotransferase	<i>glcR</i>	3739	transcriptional repressor involved in the expression of the phosphotransferase system
<i>yhaV</i>	1058	coproporphyrinogen III oxidase	<i>yitM</i>	1173	phospho-adenylsulfate sulfotransferase	<i>glcT</i>	1456	transcriptional antiterminator essential for the expression of the <i>ptsGHI</i> operon
<i>yhcB</i>	979	flavodoxin	<i>yitN</i>	1173	phospho-adenylsulfate sulfotransferase	<i>glrR</i>	1877	transcriptional repressor of the glutamine synthetase gene (<i>glrA</i>)
<i>yhfU</i>	1112	biotin biosynthesis	<i>yitO</i>	1173	phospho-adenylsulfate sulfotransferase	<i>glpP</i>	1001	transcriptional antiterminator and control of mRNA stability of <i>glpD</i>
<i>yhxA</i>	1000	adenosylmethionine-8-amino-7-oxononanoate aminotransferase	<i>yitP</i>	1173	phospho-adenylsulfate sulfotransferase	<i>glpC</i>	2014	transcriptional activator of the glutamate synthase operon (<i>gluAB</i>)
			<i>recF</i>	3	DNA repair and genetic recombination	<i>glrR</i>	2725	transcriptional repressor of the glutamate synthase operon (<i>gluAB</i>)
			<i>recN</i>	2522	DNA repair and genetic recombination	<i>gntR</i>	4113	transcriptional repressor of the gluconate operon (<i>gntRKPZ</i>)
			<i>recQ</i>	2408	ATP-dependent DNA helicase			

<i>gutR</i>	667	transcriptional activator of the sorbitol dehydrogenase gene (<i>gutA</i>)	<i>ydeC</i>	562	transcriptional regulator (AraC/XylS family)	III.5.4	TERMINATION	4
<i>hpr</i>	1073	transcriptional repressor of sporulation and extracellular proteases genes (<i>aprE</i> , <i>nprE</i> , <i>sin</i>)	<i>ydeE</i>	564	transcriptional regulator (AraC/XylS family)	<i>nusA</i>	1732	transcription termination
<i>hrcA</i>	2629	transcriptional repressor of class I heat-shock genes (<i>dnaK</i> , <i>groESL</i>)	<i>ydeF</i>	571	transcriptional regulator (GntR family) / amino-transferase (MocR-like)	<i>nusG</i>	118	transcription antitermination factor
<i>hutP</i>	4040	transcriptional activator of the histidine utilization operon (<i>hutPHUGIM</i>)	<i>ydeL</i>	574	transcriptional regulator (GntR family) / amino-transferase (MocR-like)	<i>yqhZ</i>	3904	transcriptional terminator Rho
<i>iolR</i>	4084	transcriptional repressor of the myo-inositol catabolism operon (<i>iolABCDGHIJ</i> / <i>iolRS</i>)	<i>ydeS</i>	578	transcriptional regulator (TetR/AcrR family)		2529	transcription termination
<i>kdgR</i>	2325	transcriptional repressor of the pectin utilization operon (<i>kdgRKA</i> T)	<i>ydeT</i>	579	transcriptional regulator (ArsR family)			
<i>lacR</i>	3509	transcriptional repressor of the β -galactosidase gene (<i>lacA</i>)	<i>ydhC</i>	630	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>levR</i>	2765	transcriptional activator of the levanase operon (<i>levDEFG</i> / <i>sacC</i>)	<i>ydhO</i>	609	transcriptional regulator (MarR family)		III.6	RNA MODIFICATION
<i>lexA</i>	1918	transcriptional repressor of the SOS regulon	<i>ydhQ</i>	613	transcriptional regulator (MarR family)	<i>cspR</i>	970	rRNA methylase homolog
<i>licR</i>	3963	transcriptional regulator (antiterminator) of the lichenan operon (<i>licBCA</i> H)	<i>ydhR</i>	616	transcriptional regulator (GntR family)	<i>deaD</i>	4016	ATP-dependent RNA helicase
<i>licT</i>	4012	transcriptional antiterminator required for substrate-dependent induction and catabolite repression of <i>bglPH</i>	<i>ydhS</i>	732	transcriptional regulator (TetR/AcrR family)	<i>miaA</i>	1866	tRNA isopentenylpyrophosphate transferase
<i>lmrA</i>	290	transcriptional repressor of the lincomycin operon (<i>lmrBA</i>)	<i>ydhT</i>	760	two-component response regulator [YdhT]	<i>queA</i>	2834	S-adenosylmethionine tRNA ribosyltransferase (queuosine biosynthesis)
<i>lrpA</i>	551	transcriptional Lrp-like regulator (repression of <i>glyA</i> transcription and KinB-dependent sporulation)	<i>ydhU</i>	765	transcriptional regulator (AraC/XylS family)	<i>mcs</i>	1665	ribonuclease III
<i>lrpB</i>	552	transcriptional Lrp-like regulator (repression of <i>glyA</i> transcription and KinB-dependent sporulation)	<i>ydhV</i>	790	transcriptional regulator (MarR family)	<i>mpaA</i>	4214	ribonuclease P (protein component)
<i>lrpC</i>	476	transcriptional regulator (Lrp/AsnC family)	<i>ydhW</i>	790	transcriptional regulator (MarR family)	<i>rph</i>	2901	ribonuclease PH
<i>lytR</i>	3662	attenuator role for <i>lytABC</i> and <i>lytR</i> expression	<i>ydhX</i>	790	transcriptional regulator (Lrp/AsnC family)	<i>tgt</i>	2833	tRNA-guanine transglycosylase (queuosine biosynthesis)
<i>lytT</i>	2956	two-component response regulator [LytS] involved in the rate of autolysis	<i>ydhY</i>	790	transcriptional regulator (Lrp/AsnC family)			
<i>msmR</i>	3096	transcriptional activator of multidrug-efflux transporter genes (<i>mrpA</i> and <i>mrpB</i>)	<i>ydhZ</i>	790	transcriptional regulator (Lrp/AsnC family)			
<i>mta</i>	3764	transcriptional activator of multidrug-efflux transporter genes (<i>mrpA</i> and <i>mrpB</i>)	<i>ydhA</i>	808	transcriptional regulator (AraC/XylS family)			
<i>mtrB</i>	2384	tryptophan operon RNA-binding attenuation protein (TRAP)	<i>ydhB</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>paiA</i>	3304	transcriptional repressor of sporulation, septation and degradative enzyme genes (<i>aprE</i> , <i>nprE</i> , <i>phoA</i> , <i>sacB</i>)	<i>ydhC</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>paiB</i>	3304	transcriptional repressor of sporulation and degradative enzyme genes	<i>ydhD</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>phoP</i>	2978	two-component response regulator [PhoR] involved in phosphate regulation (<i>phoA</i> , <i>phoB</i> , <i>phoD</i> , <i>resABCD</i>)	<i>ydhE</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>pkasA</i>	1781	transcriptional regulator of the polyketide synthase operon (<i>pkas</i>)	<i>ydhF</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>purR</i>	54	transcriptional repressor of the purine operon (<i>purEBCDQFMNH</i>)	<i>ydhG</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>pyrR</i>	1618	transcriptional attenuation of the pyrimidine operon (<i>pyrPBCAD</i> FE) / uracil phosphoribosyltransferase activity (minor) [pyrimidine biosynthesis]	<i>ydhH</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>rbpR</i>	3700	transcriptional repressor of the ribose operon (<i>rbpR</i> / <i>resABCD</i>)	<i>ydhI</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>resD</i>	2417	two-component response regulator [ResE] involved in aerobic and anaerobic respiration	<i>ydhJ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>ribR</i>	3001	transcriptional regulator of riboflavin biosynthesis genes	<i>ydhK</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>rocR</i>	4145	transcriptional activator of arginine utilization operons (<i>rocABC</i> , <i>rocDEF</i>)	<i>ydhL</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>sacT</i>	3906	transcriptional antiterminator involved in positive regulation of <i>sacA</i> and <i>sacP</i>	<i>ydhM</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>sacV</i>	532	transcriptional regulator of the levansucrase gene (<i>sacB</i>)	<i>ydhN</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>sacY</i>	3942	transcriptional antiterminator involved in positive regulation of levansucrase and sucrose synthesis	<i>ydhO</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>senS</i>	959	transcriptional regulator of extracellular enzyme genes (<i>amyE</i> , <i>aprE</i> , <i>nprE</i>)	<i>ydhP</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>sinR</i>	2552	transcriptional regulator of post-exponential-phase responses genes (<i>aprE</i> , <i>comK</i> , <i>kinB</i> , <i>sigD</i> , <i>spo0A</i> , <i>spo0I</i> , <i>spo0J</i> , <i>spo0K</i>)	<i>ydhQ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>slr</i>	3529	transcriptional activator of competence development and sporulation genes	<i>ydhR</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>spIA</i>	1461	transcriptional regulator of the spore photoproduct lyase operon (<i>spIA</i>)	<i>ydhS</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>spo0A</i>	2518	two-component response regulator [KinC] central for the initiation of sporulation (<i>spo0A</i> , <i>abrB</i> , <i>kinA</i> , <i>kinC</i> , <i>spo0A</i> , <i>spo0J</i> , <i>spo0K</i>) (part of phosphorelay: Spo0B-P \rightarrow Spo0A-P)	<i>ydhT</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>spo0F</i>	3809	two-component response regulator [KinA, KinB] involved in the initiation of sporulation (part of phosphorelay: Spo0F-P \rightarrow Spo0B-P)	<i>ydhU</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>spoIID</i>	3748	transcriptional regulator of σ^E - and σ^D -dependent genes	<i>ydhV</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>spoVT</i>	64	transcriptional positive and negative regulator of σ^E -dependent genes	<i>ydhW</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>tenA</i>	1242	transcriptional regulator of extracellular enzyme genes (<i>aprE</i> , <i>nprE</i> , <i>phoA</i> , <i>sacB</i>)	<i>ydhX</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>tenI</i>	1243	transcriptional activator of extracellular enzyme genes	<i>ydhY</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>tnrA</i>	1397	transcriptional pleiotropic regulator involved in global nitrogen regulation (expression of <i>nrgAB</i> , <i>nasB</i> , <i>gabP</i> , <i>ureABC</i> , <i>glnRA</i>)	<i>ydhZ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>treR</i>	853	transcriptional repressor of the trehalose operon (<i>trePAR</i>)	<i>ydhA</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>xre</i>	1321	transcriptional repressor of PBX genes	<i>ydhB</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>xylR</i>	1891	transcriptional repressor of the xyllose operon (<i>xylAB</i>)	<i>ydhC</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yacF</i>	88	transcriptional regulator (nitrogen regulation protein)	<i>ydhD</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>ybbB</i>	185	transcriptional regulator (AraC/XylS family)	<i>ydhE</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>ybdI</i>	221	two-component response regulator [YbdK]	<i>ydhF</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>ybfI</i>	244	transcriptional regulator (AraC/XylS family)	<i>ydhG</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>ybpP</i>	251	transcriptional regulator (AraC/XylS family)	<i>ydhH</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>ybgA</i>	258	transcriptional regulator (GntR family)	<i>ydhI</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>ycbB</i>	267	two-component response regulator [YcbA]	<i>ydhJ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>ycbG</i>	273	transcriptional regulator (GntR family)	<i>ydhK</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>ycbL</i>	278	two-component response regulator [YcbM]	<i>ydhL</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccH</i>	296	two-component response regulator [YccG]	<i>ydhM</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccK</i>	320	transcriptional regulator (ArsR family)	<i>ydhN</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccK</i>	341	transcriptional regulator (LysR family)	<i>ydhO</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccL</i>	412	transcriptional regulator (LysR family)	<i>ydhP</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccL</i>	426	two-component response regulator [YccI]	<i>ydhQ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccN</i>	438	transcriptional regulator (TetR/AcrR family)	<i>ydhR</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccN</i>	441	transcriptional regulator (GntR family) / amino-transferase (MocR-like)	<i>ydhS</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccN</i>	449	transcriptional regulator (DeoR family)	<i>ydhT</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccO</i>	461	transcriptional regulator (LysR family)	<i>ydhU</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccD</i>	406	transcriptional regulator (GntR family) / amino-transferase (MocR-like)	<i>ydhV</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccG</i>	439	transcriptional regulator (ArsR family)	<i>ydhW</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccG</i>	467	transcriptional antiterminator (BglG family)	<i>ydhX</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccG</i>	499	two-component response regulator [YccF]	<i>ydhY</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
<i>yccN</i>	531	transcriptional regulator (phage-related) (Xre family)	<i>ydhZ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhA</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhB</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhC</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhD</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhE</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhF</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhG</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhH</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhI</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhJ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhK</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhL</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhM</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhN</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhO</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhP</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhQ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhR</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhS</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhT</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhU</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhV</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhW</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhX</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhY</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhZ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhA</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhB</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhC</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhD</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhE</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhF</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhG</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhH</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhI</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhJ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhK</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhL</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhM</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhN</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhO</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhP</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhQ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhR</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhS</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhT</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhU</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhV</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhW</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhX</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhY</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhZ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhA</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhB</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhC</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhD</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhE</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhF</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhG</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhH</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhI</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhJ</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhK</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			
			<i>ydhL</i>	808	transcriptional regulator (GntR family) / amino-transferase (MocR-like)			

<i>lepA</i>	2632	GTP-binding protein	<i>yvE</i>	3515	spore coat polysaccharide biosynthesis	<i>xxdC</i>	1322	PBSX prophage
<i>tsf</i>	1718	elongation factor Ts	<i>yvB</i>	3384	serine protease Do	<i>xxdD</i>	1323	PBSX prophage
<i>tufA</i>	1331	elongation factor Tu	<i>ywqC</i>	3732	capsular polysaccharide biosynthesis	<i>xxdE</i>	1327	PBSX prophage
<i>ylaG</i>	1546	GTP-binding elongation factor	<i>ywqD</i>	3732	capsular polysaccharide biosynthesis	<i>xxdF</i>	1328	PBSX prophage
			<i>ywqE</i>	3731	capsular polysaccharide biosynthesis	<i>xxdG</i>	1329	PBSX prophage
III.7.5	TERMINATION3	<i>ywsC</i>	3700	capsular polyglutamate biosynthesis	<i>xxdH</i>	1330	PBSX prophage
<i>frt</i>	1720	ribosome recycling factor	<i>ywtA</i>	3698	capsular polyglutamate biosynthesis	<i>xxdI</i>	1331	PBSX prophage
<i>prfA</i>	3797	peptide chain release factor 1	<i>ywtB</i>	3698	capsular polyglutamate biosynthesis	<i>xxdJ</i>	1331	PBSX prophage
<i>prfB</i>	3627	peptide chain release factor 2	<i>yyxA</i>	4148	serine protease Do	<i>xxdK</i>	1332	PBSX prophage
						<i>xxdM</i>	1333	PBSX prophage
III.8	PROTEIN MODIFICATION27	IV.2	DETOXIFICATION68	<i>xxdN</i>	1334	PBSX prophage
<i>amhX</i>	325	amidohydrolase	<i>aadK</i>	2736	aminoglycoside 6-adenyltransferase	<i>xxdO</i>	1334	PBSX prophage
<i>lgt</i>	3593	prolipoprotein diacylglycerol transferase (lipoprotein biosynthesis)	<i>ahpC</i>	4118	alkyl hydroperoxide reductase (small subunit)	<i>xxdP</i>	1338	PBSX prophage
			<i>ahpF</i>	4119	alkyl hydroperoxide reductase (large subunit)	<i>xxdQ</i>	1339	PBSX prophage
<i>map</i>	147	methionine aminopeptidase	<i>bmrU</i>	2493	NADH dehydrogenase	<i>xxdR</i>	1340	PBSX prophage
<i>pcp</i>	287	pyrrolidone-carboxylate peptidase				<i>xxdS</i>	1340	PBSX prophage
<i>ppbB</i>	2435	peptidyl-prolyl isomerase				<i>xxdT</i>	1341	PBSX prophage
<i>prkA</i>	973	serine protein kinase	<i>cah</i>	342	cephalosporin C deacetylase	<i>xxdU</i>	1342	PBSX prophage
<i>tgl</i>	3212	transglutaminase	<i>cypA</i>	2732	cytochrome P450-like enzyme	<i>xxdV</i>	1343	PBSX prophage
<i>ybdM</i>	224	protein kinase	<i>cypX</i>	3603	cytochrome P450-like enzyme	<i>xxdW</i>	1345	PBSX prophage
<i>ydiC</i>	642	glycoprotein endopeptidase	<i>katA</i>	960	vegetative catalase 1	<i>xxdX</i>	1345	PBSX prophage
<i>ydiD</i>	643	ribosomal-protein-alanine N-acetyltransferase	<i>katB</i>	4009	catalase 2	<i>xxdY</i>	1345	PBSX prophage lytic exoenzyme
<i>ydiE</i>	643	glycoprotein endopeptidase	<i>katX</i>	3964	catalase	<i>xtmA</i>	1325	PBSX terminase (small subunit)
<i>ytkJ</i>	862	protein-tyrosine phosphatase	<i>ksgA</i>	51	dimethyladenosine transferase (kasugamycin resistance)	<i>xtmB</i>	1325	PBSX terminase (large subunit)
<i>ytlG</i>	840	methionine aminopeptidase				<i>xtrA</i>	1324	PBSX prophage
<i>yvCk</i>	1261	ribosomal-protein-alanine N-acetyltransferase	<i>mmr</i>	3657	methylerythromycin A resistance protein	<i>ycdD</i>	304	L-alanyl-D-glutamate peptidase
<i>yvB</i>	1528	formylmethionine deformylase	<i>padC</i>	3532	ferulate decarboxylase	<i>ydcL</i>	530	integrase
<i>yvY</i>	1453	Xaa-Pro dipeptidase	<i>penP</i>	2048	β -lactamase	<i>ydcM</i>	531	immunity region protein in prophage
<i>yvP</i>	1651	protein kinase	<i>pkS</i>	1859	hydrolyase of the polyketide produced by the	<i>yhgE</i>	1090	phage infection protein
<i>yppP</i>	2287	peptide methionine sulfoxide reductase				<i>yjbl</i>	1235	lytic transglycosylase
<i>yqeT</i>	2624	ribosomal protein L11 methyltransferase	<i>sodA</i>	2585	superoxide dismutase	<i>yjgB</i>	1318	phage-related replication protein
<i>yqhT</i>	2539	Xaa-Pro dipeptidase	<i>sodF</i>	2103	superoxide dismutase	<i>ymaC</i>	1863	phage-related protein
<i>ytl</i>	3020	protease IV	<i>tetL</i>	4188	tetracycline resistance leader peptide	<i>ymaH</i>	1867	host factor-1 protein
<i>ytlP</i>	3068	Xaa-His dipeptidase	<i>thdF</i>	4212	thiophen and furan oxidation	<i>ymlD</i>	1755	phage-related protein
<i>yvA</i>	3105	protein kinase	<i>tmmB</i>	339	unicarmycin resistance	<i>ymlE</i>	1756	phage-related protein
<i>yvM</i>	3150	prolyl aminopeptidase	<i>yaaN</i>	36	toxic cation resistance	<i>ymdL</i>	1914	phage-related replication protein
<i>yvE</i>	3297	leucyl aminopeptidase	<i>ydbE</i>	190	β -lactamase	<i>yobD</i>	2075	phage-related pre-neck appendage protein
<i>yvE</i>	3791	protein-tyrosine phosphatase	<i>ybfO</i>	205	erythromycin esterase	<i>yokA</i>	2284	DNA recombinase
<i>yyaL</i>	4102	serine/threonine protein kinase	<i>ybl</i>	229	β -lactamase	<i>yokL</i>	2274	phage-related protein
			<i>ycbI</i>	276	viomycin phosphotransferase	<i>yolB</i>	2272	phage-related protein
III.9	PROTEIN FOLDING8	<i>ycbR</i>	283	toxic cation resistance protein	<i>yomA</i>	2264	holin
<i>dnaK</i>	2627	class I heat-shock protein (chaperonin)	<i>yceC</i>	312	tellurium resistance protein	<i>yomJ</i>	2248	phage-related immunity protein
<i>groEL</i>	650	class I heat-shock protein (chaperonin)	<i>yceD</i>	312	tellurium resistance protein	<i>yomP</i>	2243	phage-related protein
<i>groES</i>	650	class I heat-shock protein (chaperonin)	<i>yceE</i>	313	tellurium resistance protein	<i>yomR</i>	2242	phage-related protein
<i>tig</i>	2887	trigger factor (prolyl isomerase)	<i>yceF</i>	314	tellurium resistance protein	<i>yomS</i>	2241	phage-related lytic exoenzyme
<i>ykkC</i>	1376	chaperonin	<i>yceH</i>	316	toxic anion resistance protein	<i>yqqD</i>	2200	phage-related DNA-binding protein anti-repressor
<i>ykkD</i>	1376	chaperonin	<i>yceF</i>	457	lactam utilization protein	<i>yqz</i>	2190	phage-related protein
<i>yvD</i>	3541	chaperonin	<i>ydbD</i>	496	manganese-containing catalase	<i>yqC</i>	2160	phage-related endonuclease
<i>yvS</i>	3541	chaperonin	<i>ydbB</i>	581	antibiotic resistance protein	<i>yqbB</i>	2700	phage-related protein
IV	OTHER FUNCTIONS	289	<i>ydhE</i>	618	macrolide glycosyltransferase	<i>yqal</i>	2696	phage-related protein
IV.1	ADAPTATION TO ATYPICAL CONDITIONS72	<i>yerP</i>	732	acriflavine resistance protein	<i>yqak</i>	2695	phage-related protein
<i>bssA</i>	2304	glutathione peroxidase	<i>yetM</i>	790	salicylate 1-monooxygenase	<i>yqam</i>	2694	phage-related protein
<i>clpC</i>	104	class III stress response-related ATPase (repressor of competence)	<i>yetO</i>	792	cytochrome P450 / NADPH-cytochrome P450 reductase	<i>yqaO</i>	2692	phage-related protein
<i>clpE</i>	1437	ATP-dependent Clp protease-like	<i>yilM</i>	836	nitric-oxide synthase	<i>yqaS</i>	2690	phage-related terminase large subunit
<i>clpP</i>	3545	ATP-dependent Clp protease proteolytic subunit (class III heat-shock protein)	<i>ylnC</i>	804	fosmidmycin resistance protein	<i>yqaT</i>	2689	phage-related terminase small subunit
<i>clpQ</i>	1688	B-type subunit of the 20S proteasome	<i>ygaF</i>	943	thiol-specific antioxidant protein	<i>yqbA</i>	2688	phage-related protein
<i>clpX</i>	2885	ATP-dependent Clp protease ATP-binding subunit (class III heat-shock protein)	<i>yhgG</i>	1122	monooxygenase	<i>yqbD</i>	2684	phage-related protein
<i>clpY</i>	1688	ATP-dependent Clp protease-like	<i>yibY</i>	1169	chloride peroxidase	<i>yqde</i>	2683	phage-related protein
<i>csbB</i>	930	stress response protein	<i>yijB</i>	1291	monooxygenase	<i>yqdeH</i>	2682	phage-related protein
<i>csbP</i>	984	major cold-shock protein	<i>yijG</i>	1292	macrolide glycosyltransferase	<i>yqbl</i>	2681	phage-related protein
<i>csbC</i>	559	cold-shock protein	<i>yikA</i>	1366	immunity to bacteriotoxins	<i>yqbl</i>	2681	phage-related protein
<i>csbD</i>	2307	cold-shock protein	<i>ykkB</i>	1375	N-acetyltransferase	<i>yqk</i>	2680	phage-related protein
<i>cstA</i>	2937	carbon starvation-induced protein	<i>ykoY</i>	1410	toxic anion resistance protein	<i>yqbl</i>	2679	phage-related protein
<i>ctc</i>	59	general stress protein	<i>yndN</i>	1916	fosfomycin resistance protein	<i>yqbM</i>	2679	phage-related protein
<i>degQ</i>	3256	degradative enzyme production	<i>yocD</i>	2088	immunity to bacteriotoxins	<i>yqbN</i>	2677	phage-related protein
<i>degR</i>	3206	degradative enzyme production	<i>yoiK</i>	2117	macrolide glycosyltransferase	<i>yqbO</i>	2677	phage-related protein
<i>dnaI</i>	2625	heat-shock protein (activation of DnaK)	<i>yoiM</i>	2115	superoxide dismutase	<i>yqbP</i>	2672	phage-related protein
<i>dps</i>	3136	stress- and starvation-induced gene controlled by	<i>yokD</i>	2281	aminoglycoside N ³ -acetyltransferase	<i>yqbQ</i>	2671	phage-related protein
			<i>yqcm</i>	2655	arsenate reductase	<i>yqbr</i>	2670	phage-related protein
<i>gbsA</i>	3186	glycine betaine aldehyde dehydrogenase (osmoprotection)	<i>yqpP</i>	2596	penicillin tolerance	<i>yqbt</i>	2670	phage-related protein
<i>gbsB</i>	3184	alcohol dehydrogenase (osmoprotection)	<i>yrlJ</i>	2776	cytochrome P450 / NADPH-cytochrome P450 reductase	<i>yqcA</i>	2669	phage-related protein
<i>griP</i>	2628	heat-shock protein (activation of DnaK)	<i>ytrB</i>	2736	2-nitropropane dioxygenase	<i>yqcC</i>	2668	phage-related protein
<i>gsiB</i>	494	general stress protein	<i>ytlG</i>	3017	thiol peroxidase	<i>yqcD</i>	2667	phage-related protein
<i>gspA</i>	3944	general stress protein	<i>ytnJ</i>	3002	nitrotriacetate monooxygenase	<i>yqcE</i>	2666	phage-related protein
<i>hit</i>	1076	Hit-like protein involved in cell-cycle regulation	<i>yubB</i>	3195	bactitracin resistance protein (undecaprenol kinase)	<i>yqxG</i>	2666	phage-related lytic exoenzyme
<i>htgP</i>	4090	class III heat-shock protein (chaperonin)				<i>yqxH</i>	2665	holin
<i>htrA</i>	1359	serine protease Do (heat-shock protein)	<i>yusi</i>	3366	arsenate reductase	IV.5	TRANSPOSON AND IS10
<i>ispU</i>	1387	activation of σ^E	<i>yvtT</i>	3487	alkanol monooxygenase	<i>ydcP</i>	533	transposon protein
<i>lonA</i>	2882	class III heat-shock ATP-dependent protease	<i>yvpP</i>	3543	reticulone oxidase	<i>ydcQ</i>	533	transposon protein
<i>lonB</i>	2884	Lon-like ATP-dependent protease	<i>ywch</i>	3810	monooxygenase	<i>ydr</i>	535	transposon protein
<i>mrqA</i>	3983	metallorepression DNA-binding stress protein	<i>ywnH</i>	3760	phosphothricin acetyltransferase	<i>ydbB</i>	537	transposon protein
<i>rsbR</i>	519	positive regulator of σ^E activity (interaction with RsbS)	<i>yxel</i>	4062	penicillin amidase	<i>ydeB</i>	538	transposon protein
<i>rsbS</i>	520	negative regulator of σ^E activity (antagonist of RsbT)	<i>yxkC</i>	4061	monooxygenase	<i>ydhH</i>	544	transposon protein
<i>rsbT</i>	520	positive regulator of σ^E activity (switch protein/serine kinase [RsbS])	<i>yysR</i>	4185	streptothricin acetyl-transferase	<i>yefB</i>	739	site-specific recombinase
<i>rsbU</i>	521	indirect positive regulator of σ^E activity (serine phosphatase [RsbV-P])	IV.3	ANTIBIOTIC PRODUCTION30	<i>yefC</i>	739	resolvase
<i>rsbV</i>	522	positive regulator of σ^E activity (anti-anti-sigma factor [RsbV])	<i>pkSB</i>	1782	involved in polyketide synthesis	<i>yneB</i>	1918	resolvase
<i>rsbW</i>	522	negative regulator of σ^E activity (switch protein/serine kinase [RsbV], anti-sigma factor [σ^E])	<i>pkSC</i>	1783	involved in polyketide synthesis	<i>yocA</i>	2085	transposon-related protein
<i>rsbX</i>	523	indirect negative regulator of σ^E activity (serine phosphatase [RsbS-P])	<i>pkSD</i>	1785	involved in polyketide synthesis	IV.6	MISCELLANEOUS26
<i>ycdH</i>	308	adhesion protein	<i>pkSE</i>	1785	involved in polyketide synthesis	<i>bex</i>	2610	GTP-binding protein
<i>ydaG</i>	473	general stress protein	<i>pkSF</i>	1788	involved in polyketide synthesis	<i>csbA</i>	3614	putative membrane protein
<i>ytlR</i>	910	surface adhesion	<i>pkSG</i>	1789	involved in polyketide synthesis	<i>csbB</i>	36	σ^E -transcribed gene
<i>yvK</i>	1414	heat-shock protein	<i>pkSH</i>	1790	involved in polyketide synthesis	<i>ctaG</i>	1564	function unknown
<i>yvZA</i>	1381	general stress protein	<i>pkSI</i>	1791	involved in polyketide synthesis	<i>eag</i>	1430	small membrane protein
<i>yvA</i>	1637	fibrinectin-binding protein	<i>pkSL</i>	1792	involved in polyketide synthesis	<i>ecsC</i>	1079	function unknown
<i>yvU</i>	1655	alkaline-shock protein	<i>pkSK</i>	1794	polyketide synthase	<i>mmgE</i>	2509	function unknown
<i>yvB</i>	1875	GTP-binding protein protease modulator	<i>pkSL</i>	1808	polyketide synthase	<i>nifZ</i>	3027	NiF protein homologue
<i>yzfF</i>	1880	δ -endotoxin	<i>pkSM</i>	1821	polyketide synthase	<i>sapB</i>	726	mutant activates alkaline phosphatase during sporulation independently of σ^E and σ^F
<i>yocK</i>	2097	general stress protein	<i>pkSN</i>	1834	polyketide synthase	<i>sbp</i>	1595	small basic protein
<i>yocM</i>	2098	small heat-shock protein	<i>pkSP</i>	1835	polyketide synthase	<i>veg</i>	53	function unknown
<i>yodU</i>	2151	capsular polysaccharide biosynthesis	<i>pkSR</i>	1850	polyketide synthase	<i>yael</i>	102	creatine kinase
<i>yokG</i>	2279	δ -endotoxin	<i>ppsA</i>	1997	peptide synthetase	<i>ybaL</i>	157	ATP-binding Mrp-like protein
<i>yqpP</i>	2286	capsular polysaccharide biosynthesis	<i>ppsB</i>	1990	peptide synthetase	<i>ybcU</i>	287	NiF protein homologue
<i>yvG</i>	3047	general stress protein	<i>ppsC</i>	1982	peptide synthetase	<i>yerN</i>	730	pet112-like protein
<i>yvJ</i>	3046	general stress protein	<i>ppsD</i>	1974	peptide synthetase	<i>yhdP</i>	1033	hemolysin
<i>yvK</i>	3529	capsular polysaccharide biosynthesis	<i>ppsE</i>	1963	peptide synthetase	<i>yhdT</i>	1035	hemolysin
<i>yvE</i>	3528	capsular polysaccharide biosynthesis	<i>sbo</i>	3835	subtilisin A	<i>yheG</i>	1049	calcium-binding protein
<i>yvM</i>	3527	capsular polysaccharide biosynthesis	<i>sfp</i>	408	surfactin production	<i>yplQ</i>	2295	hemolysin III homologue
<i>yvN</i>	3525	capsular polysaccharide biosynthesis	<i>srfAA</i>	377	surfactin synthetase / competence	<i>yqxG</i>	2523	hemolysin-like
<i>yvO</i>	3524	exopolysaccharide biosynthesis	<i>srfAB</i>	387	surfactin synthetase / competence	<i>yvA</i>	2720	hemolysin-like
<i>yvP</i>	3523	capsular polysaccharide biosynthesis	<i>srfAC</i>	398	surfactin synthetase / competence	<i>yrvO</i>	2811	NiF protein homologue
<i>yvQ</i>	3522	capsular polysaccharide biosynthesis	<i>srfAD</i>	402	surfactin synthetase / competence	<i>yuaG</i>	3187	epidermal surface antigen
<i>yvR</i>	3521	spore coat polysaccharide biosynthesis	<i>sunA</i>	2269	subtilisin 1681 antibiotic antimicrobial precursor peptide	<i>yurV</i>	3357	NiF protein homologue
<i>yvT</i>	3519	capsular polysaccharide biosynthesis	<i>yomB</i>	2264	bacteriocin	<i>yurW</i>	3358	NiF protein homologue
<i>yvC</i>	3517	capsular polysaccharide biosynthesis	<i>yukL</i>	3282	antibiotic synthetase	<i>yutI</i>	3309	NiF protein homologue
			<i>yukM</i>	3283	antibiotic synthetase	V	SIMILAR TO UNKNOWN PROTEINS	668
IV.4	PHAGE-RELATED FUNCTIONS83	IV.4	PHAGE-RELATED FUNCTIONS83	V.1	FROM <i>B. SUBTILIS</i>177
<i>codV</i>	1687	integrase/recombinase	<i>ripX</i>	2449	integrase/recombinase	V.2	FROM OTHER ORGANISMS491
<i>xhIA</i>	1346	involved in cell lysis upon induction of PBSX	<i>xhIB</i>	1346	hydrolysis of 5-bromo 4-chloroindolyl phosphate upon induction of PBSX (holin)	VI	NO SIMILARITY	1053
<i>xxdA</i>	1320	PBSX prophage						
<i>xxdB</i>	1321	PBSX prophage						